

GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:46 ; Search time 225.158 Seconds  
(without alignments) 3231.380 Million cell updates/sec

Title: US-09-787-562-1  
Perfect score: 25  
Sequence: 1 cgcgtcggtgcaggacatgacaaat 25

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
                  Maximum Match 10%
                  Listing first 45

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Database : GenEmbl : \*

1: gp\_ba:\*  
2: gb\_htg:\*  
3: gb\_in:\*  
4: gb\_om:\*  
5: gb\_ov:\*  
6: gb\_pat:\*  
7: gb\_pl:\*  
8: gb\_pl1:\*  
9: gb\_pr:\*  
10: gb\_ro:\*  
11: gb\_sts:\*  
12: gb\_sy:\*  
13: gb\_un:\*  
14: gb\_vt:\*  
15: en\_ba:\*  
16: en\_fun:\*  
17: em\_hum:\*  
18: em\_in:\*  
19: em\_mu:\*  
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21: em\_ov:\*  
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28: en\_un:\*  
29: em\_vt:\*  
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31: em\_htg\_inv:\*  
32: em\_htg\_other:\*  
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35: em\_htg\_rod:\*  
36: em\_htg\_mam:\*  
37: em\_htg\_vrt:\*  
38: em\_sy:\*  
39: em\_tgo\_hum:\*  
40: em\_tgo\_mus:\*  
41: em\_tgo\_other:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query %		DB	ID	Description
		Match	Length			
1	25	100.0	25	6	AX023659	Sequence
2	19.2	76.8	273785	1	SME591793	AL591793 Sinorhizo
3	19.2	76.8	338579	1	AP003004	AX023660 Sequence
4	19	76.0	19	6	AX023660	AX023660 Sequence
5	18.6	74.4	695	8	HOS271030	AJ271030 Hansenias
6	18.6	74.4	704	8	HVS271033	AJ271033 Hansenias
7	18.6	74.4	3670	1	VFU01417	U01417 Vibrio furn
8	18.6	74.4	3670	6	AR022345	AR022345 Sequence
9	18.6	74.4	3670	6	AR086949	AR086949 Sequence
10	18.6	74.4	217812	2	AC130788	AC130788 Bos tauri
11	18.4	73.6	117962	2	AP003914	AP003914 Oriza sat
12	18.4	73.6	154255	2	AP004632	AP004632 Oriza sat
13	18	72.0	24	6	AX023673	AX023673 Sequence
14	18	72.0	24	6	AX048713	AX048713 Sequence
15	18	72.0	24	6	AX048713	AX048713 Sequence
16	18	72.0	366	5	AF353996	AF353996 Cyprinus
17	18	72.0	513	6	AX150246	AX150246 Sequence
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24	18	72.0	4768	6	AX352704	AX352704 Sequence
25	18	72.0	4847	6	AX114854	AX114854 Sequence
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31	18	72.0	5608	12	AF092169	AF092169 Cloning v
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36	18	72.0	5608	12	AF092542	AF092542 Cloning v
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## ALIGNMENTS

RESULT 1	AX023659	Sequence 1 from Patent WO01/7371.	25 bp	DNA	linear	PAT 15-SEP-2000
LOCUS	AX023659					
DEFINITION	AX023659					
ACCESSION	AX023659					
VERSION	AX023659.1	GI:10184020				
KEYWORDS						
SOURCE						
ORGANISM						
		Mus sp.				
		Mus sp.				
		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
		Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
REFERENCE		1 (bases 1 to 25)				
AUTHORS		Binley, K.M. and Naylor, S.				
TITLE		Polynucleotide constructs and uses thereof				
JOURNAL		Patent: WO 001371-A 1 30-MAR-2000;				

Mon Jan 6 15:20:16 2003

BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD (GB)

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Location/Qualifiers  
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Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 CGCGTCGGTCGACGACGACAAAT 25  
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Db 1 CGCGTCGGTCGACGACGACAAAT 25  
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LOCUS Sinorhizobium meliloti 1021 complete chromosome; segment 12/12.  
DEFINITION  
ACCESSION AL591793 AL591688  
VERSION AL591793.1 GI:15076142  
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SOURCE Sinorhizobium meliloti.  
ORGANISM Sinorhizobium meliloti  
Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group; Rhizobiaceae; Sinorhizobium.

REFERENCE  
AUTHORS  
1 (bases 1 to 273785)  
Capela,D., Barloy-Hubier,F., Gouzy,J., Bothe,G., Ampe,F., Batut,J., Boistard,P., Becker,A., Boutry,M., Cadieu,E., Dreano,S., Gloux,S., Godrie,T., Goffeau,A., Kahn,D., Kiss,E., Lelaure,V., Masuy,D., Pohl,T., Portetelle,D., Puhler,A., Purnelle,B., Ramsperger,U., Renard,C., Thebault,P., Vandenbol,M., Weidner,S. and Galibert,F.  
Analysis of the chromosome sequence of the legume symbiont Sinorhizobium meliloti strain 1021  
Proc. Natl. Acad. Sci. U.S.A. 98 (17), 9877-9882 (2001)  
21396507  
11481430

TITLE  
JOURNAL  
MEDLINE  
PUBMED  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT  
Submitted (26-JUL-2001) Gouzy J., Submitted on behalf of the MELILO EU Consortium  
MELILO EU Consortium:  
Laboratoire de Biologie Moleculaire des Relations  
Plantes-Microorganismes, UMR215-CNRS-INRA, BP27, F-31326 Castanet,  
France, Laboratoire de Genetique et Developpement UMR6061-CNRS,  
Faculte de Medecine, 2 avenue du Pr. Leon Bernard, F-35043 Rennes,  
France, CATC GmbH, Fritz-Arnold-str. 23, D-78467 Konstanz, Germany,  
Universitaet Bielefeld, Biologie IV (Genetik) Universitaetstr 25,  
D-33615 Bielefeld, Germany, Unite de Biochimie physiologique,  
Universite Catholique de Louvain, Place Croix du Sud 2, Bte 20,  
B-1348 Louvain-la-Neuve, Belgium, Unite de Microbiologie, Faculte  
des Sciences Agronomiques de Gembloux, Avenue Marechal Juin 6,  
B-5030 Gembloux, Belgium. E-mail:Jerome.Gouzy@toulouse.inra.fr  
http://sequence.toulouse.inra.fr/meliloti.html.

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LOCUS Mesorhizobium loti DNA, complete genome, section 11/21.  
DEFINITION AP003004 BA000012.  
ACCESSION AP003004.2 GI:14024426  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Mesorhizobium loti (strain:MAFF303099) DNA.  
Mesorhizobium loti  
Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;  
Phyllobacteriaceae; Mesorhizobium.  
REFERENCE  
1 (sites)  
AUTHORS  
Kaneko, T., Nakamura, Y., Sato, S., Asamizu, E., Kato, T., Sasamoto, S.,  
Watanabe, A., Iidesawa, K., Ishikawa, A., Kawashima, K., Kimura, T.,  
Kishida, Y., Kiyokawa, C., Kohara, M., Matsumoto, M., Matsuno, A.,  
Mochizuki, Y., Nakayama, S., Nakazaki, N., Shimpo, S., Sugimoto, M.,  
Takeuchi, C., Yamada, M. and Tabata, S.  
Complete genome structure of the nitrogen-fixing symbiotic  
bacterium Mesorhizobium loti  
DNA Res. 7 (6), 331-338 (2000)  
21082930  
REFERENCE  
2 (bases 1 to 338579)  
AUTHORS  
Kaneko, T.  
DIRECT SUBMISSION  
SUBMITTED (05-DEC-2000) Takakazu Kaneko, Kazusa DNA Research  
Institute, The First Laboratory for Plant Gene Research; Yana  
1532-3, Kisarazu, Chiba 292-0812, Japan  
(E-mail:kaneko@kazusa.or.jp/rhizobase/  
URL:http://www.kazusa.or.jp/rhizobase/  
Tel:81-438-52-3935(ex.2338), Fax:81-438-52-3934)  
On May 11, 2001 this sequence version replaced gi:11994979.  
COMMENT  
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AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE Binley.K.M. and Naylor.S.
JOURNAL Polynucleotide constructs and uses thereof
PATENT: WO 0017371-A 2 30-MAR-2000;
BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD
(GB)

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VERSION AJ271030.1 GI:11691808
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ORGANISM Hanseniaspora osmophila
Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
Saccharomycetales; Saccharomycodaceae; Hanseniaspora.
1 (bases 1 to 695)
AUTHORS Esteve-Zarzoso,B., Peris-Toran,M., Ramon,D. and Querol,A.
TITLE Molecular characterisation of the species to the genus
Hanseniaspora
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 695)
AUTHORS Esteve-Zarzoso,B.
TITLE Direct Submission
JOURNAL Submitted (10-JAN-2000) Esteve-Zarzoso B., Departament de
Biotechnologia, Institut d'Agroquímica i Tecnologia d'Aliments,
CSIC, P.O. Box 73, 46100 Burjassot, Valencia, SPAIN

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ORGANISM Vibrio furnissii  
Bacteria; Proteobacteria; gamma subdivision; Vibrionaceae; Vibrio.  
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Keyhani.N.O. and Roseman.S.  
The chitin catabolic cascade in the marine bacterium Vibrio furnissii. Molecular cloning, isolation, and characterization of a periplasmic beta-N-acetylglucosaminidase  
J Biol Chem. 271 (52), 33425-33432 (1996)  
97125983  
8969205  
2 (bases 1 to 3670)  
Keyhani.N.O. and Roseman.S.  
Direct Submission  
Submitted (27-NOV-1995) Nemat O. Keyhani, Biology, Johns Hopkins University, 3400 N. Charles St., Baltimore, MD 21218, USA

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SOURCE Vibrio furnissii.  
ORGANISM Vibrio furnissii  
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1 (bases 1 to 3670)  
Keyhani.N.O. and Roseman.S.  
The chitin catabolic cascade in the marine bacterium Vibrio furnissii. Molecular cloning, isolation, and characterization of a periplasmic beta-N-acetylglucosaminidase  
J Biol Chem. 271 (52), 33425-33432 (1996)  
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8969205  
2 (bases 1 to 3670)  
Keyhani.N.O. and Roseman.S.  
Direct Submission  
Submitted (27-NOV-1995) Nemat O. Keyhani, Biology, Johns Hopkins University, 3400 N. Charles St., Baltimore, MD 21218, USA

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/translation="MNYRIDFAVLSEHPQCRGLLHNLSDQDLKAWSLHFTIDRYI QPDSISRHQIVGVSFCSLTPEQDVINSNSHYCEFSIKTAPFFHYITDGIKAFFVQ INDVEPRHDVITPITALASPYRSEIPATDAATLSLPKPNHIERLDGEFALTAG SOISLOSSCAETAATWLKQELTHLYOWPHDIGSADIVLTNPNTLDGAYLLISVDRKP IRLEASSHIGFVASATLLQIVRPDGNLAVPHIVIKADAPFRYRGMLDLCARHPPL ERKRLINLQAHYKFTNFHMLTDDEGWREIKSLPQLTDIGAWRGVDEVLPEOYSLL TERKGYTQEEIHEVITAYAAERGITVPEIDIPGHSRAAKALPFWLDFEDDSQYR SIQYINDVLSPALPGYRFLDCVLEEVAAFPFSHTIHGADEVDPGVGWSKPCQAL MAESGYDAKELQGHLLRYAEKLLSKRGWMEEAQHGDKVSKDTVIYSLVSEQAA LNCARQGFVDLQPGQFTYLDIAQYAPPEPGVDWAGVTPLERAYRVEPLVEVPEHDP LRKRLIGQALWCELNNQDRMDYMLYPLRTALAGSLDTKIPA"

BASE COUNT 858 a 1046 c 958 g 808 t  
ORIGIN

Query Match 74.4%; Score 18.6; DB 1; Length 3670;  
Best Local Similarity 84.0%; Pred. No. 1.1e+03;  
Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Y 1 CGCGTCGGTCAGGACGTGACAAAT 25  
||||| ||||||| || ||| |||  
b 542 CGCGTCGTGCAGGATGTAACCAAT 566

RESULT 7  
VFU41417 3670 bp DNA linear BCT 24-DEC-1996  
LOCUS Vibrio furnissii beta-N-acetylhexosaminidase gene, complete cds.  
DEFINITION U41417  
ACCESSION U41417.1 GI:1698441  
VERSION  
KEYWORDS  
SOURCE Vibrio furnissii.  
ORGANISM Vibrio furnissii  
Bacteria; Proteobacteria; gamma subdivision; Vibrionaceae; Vibrio.  
1 (bases 1 to 3670)  
Keyhani.N.O. and Roseman.S.  
The chitin catabolic cascade in the marine bacterium Vibrio furnissii. Molecular cloning, isolation, and characterization of a periplasmic beta-N-acetylglucosaminidase  
J Biol Chem. 271 (52), 33425-33432 (1996)  
97125983  
8969205  
2 (bases 1 to 3670)  
Keyhani.N.O. and Roseman.S.  
Direct Submission  
Submitted (27-NOV-1995) Nemat O. Keyhani, Biology, Johns Hopkins University, 3400 N. Charles St., Baltimore, MD 21218, USA

FEATURES  
source location/Qualifiers  
1..3670  
/organism="Vibrio furnissii"  
/db\_xref="taxon:29494"  
845..2680  
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BASE COUNT 858 a 1046 c 958 g 808 t  
ORIGIN

Query Match 74.4%; Score 18.6; DB 1; Length 3670;  
Best Local Similarity 84.0%; Pred. No. 1.1e+03;  
Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Y 1 CGCGTCGGTCAGGACGTGACAAAT 25  
||||| ||||||| || ||| |||  
b 542 CGCGTCGTGCAGGATGTAACCAAT 566

RESULT 7  
VFU41417 3670 bp DNA linear BCT 24-DEC-1996  
LOCUS Vibrio furnissii beta-N-acetylhexosaminidase gene, complete cds.  
DEFINITION U41417  
ACCESSION U41417.1 GI:1698441  
VERSION  
KEYWORDS  
SOURCE Vibrio furnissii.  
ORGANISM Vibrio furnissii  
Bacteria; Proteobacteria; gamma subdivision; Vibrionaceae; Vibrio.  
1 (bases 1 to 3670)  
Keyhani.N.O. and Roseman.S.  
The chitin catabolic cascade in the marine bacterium Vibrio furnissii. Molecular cloning, isolation, and characterization of a periplasmic beta-N-acetylglucosaminidase  
J Biol Chem. 271 (52), 33425-33432 (1996)  
97125983  
8969205  
2 (bases 1 to 3670)  
Keyhani.N.O. and Roseman.S.  
Direct Submission  
Submitted (27-NOV-1995) Nemat O. Keyhani, Biology, Johns Hopkins University, 3400 N. Charles St., Baltimore, MD 21218, USA

FEATURES  
source location/Qualifiers  
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/organism="Vibrio furnissii"  
/db\_xref="taxon:29494"  
845..2680  
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/translation="MNYRIDFAVLSEHPQCRGLLHNLSDQDLKAWSLHFTIDRYI QPDSISRHQIVGVSFCSLTPEQDVINSNSHYCEFSIKTAPFFHYITDGIKAFFVQ INDVEPRHDVITPITALASPYRSEIPATDAATLSLPKPNHIERLDGEFALTAG SOISLOSSCAETAATWLKQELTHLYOWPHDIGSADIVLTNPNTLDGAYLLISVDRKP IRLEASSHIGFVASATLLQIVRPDGNLAVPHIVIKADAPFRYRGMLDLCARHPPL ERKRLINLQAHYKFTNFHMLTDDEGWREIKSLPQLTDIGAWRGVDEVLPEOYSLL TERKGYTQEEIHEVITAYAAERGITVPEIDIPGHSRAAKALPFWLDFEDDSQYR SIQYINDVLSPALPGYRFLDCVLEEVAAFPFSHTIHGADEVDPGVGWSKPCQAL MAESGYDAKELQGHLLRYAEKLLSKRGWMEEAQHGDKVSKDTVIYSLVSEQAA LNCARQGFVDLQPGQFTYLDIAQYAPPEPGVDWAGVTPLERAYRVEPLVEVPEHDP LRKRLIGQALWCELNNQDRMDYMLYPLRTALAGSLDTKIPA"

BASE COUNT 858 a 1046 c 958 g 808 t  
ORIGIN

Query Match 74.4%; Score 18.6; DB 1; Length 3670;  
Best Local Similarity 84.0%; Pred. No. 1.1e+03;  
Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Y 1 CGCGTCGGTCAGGACGTGACAAAT 25  
||||| ||||||| || ||| |||  
b 542 CGCGTCGTGCAGGATGTAACCAAT 566

RESULT 7  
VFU41417 3670 bp DNA linear BCT 24-DEC-1996  
LOCUS Vibrio furn

Thomas,P.J., Touchman,J.W., Tsurgeon,C., Vogt,J.L., Walker,M.A.,  
 Wetherby,K.D., Wiggins,L., Young,A., Zhang,L.-H. and Green,E.D.  
 NISC Comparative Sequencing Initiative  
 Unpublished  
 2 (bases 1 to 217812)  
 Green,E.D.

TITLE  
 JOURNAL  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL

Direct Submission  
 Submitted (14-AUG-2002) NIH Intramural Sequencing Center, 8717  
 Government Circle, Gaithersburg, MD 20877, USA  
 ----- Genome Center  
 Center: NIH Intramural Sequencing Center  
 Center code: NISC  
 Web site: <http://www.nisc.nih.gov>  
 Contact: [nisc\\_zoo@nhri.nih.gov](mailto:nisc_zoo@nhri.nih.gov)  
 ----- Project Information  
 Center project name: crz  
 Center clone name: 331024  
 ----- Summary Statistics

COMMENT

Sequencing vector: plasmid; n/a; 100% of reads  
 Chemistry: Dye-terminator Big Dye; 100% of reads  
 Assembly program: Phrap; version 0.990319  
 Consensus quality: 214289 bases at least Q40  
 Consensus quality: 215318 bases at least Q30  
 Consensus quality: 215867 bases at least Q20  
 Insert size: 208000; agarose-fp  
 Insert size: 217412; sum-of-contigs  
 Quality coverage: 8.46x in Q20 bases; agarose-fp  
 Quality coverage: 8.10x in Q20 bases; sum-of-contigs  
 -----

\* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 5 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

\* 1 3022: contig of 3022 bp in length  
 \* 3023 3122: gap of unknown length  
 \* 3123 36890: contig of 33768 bp in length  
 \* 36891 36930: gap of unknown length  
 \* 36931 66601: contig of 29611 bp in length  
 \* 66602 66701: gap of unknown length  
 \* 66702 120837: contig of 54136 bp in length  
 \* 120838 120937: gap of unknown length  
 \* 120938 217812: contig of 96875 bp in length.

FEATURES  
 source

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 /db\_xref="taxon:9913"  
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 /clone\_lib="RP42"  
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 66702. .120837  
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 120938. .217812  
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misc\_feature

BASE COUNT 51110 a 58399 c 58136 g 49765 t 402 others  
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 Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CGCGTCGGTCGACGACCTGACAAAT 25  
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 Db 150195 CGCGTCGTGTGCACACCTGACAAAT 150171

RESULT 11  
 AP003914/c  
 LOCUS  
 DEFINITION

ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

The nucleotide sequence of this BAC clone was generated by  
 combining Monsanto and RGP-Japan sequencing data.  
 NOTE: It currently consists of 1 contigs. Gaps between the contigs  
 are represented as runs of N. The order of the pieces is believed  
 to be correct as given, however the sizes of the gaps between them  
 are based on estimates that have provided by the submitter. This  
 sequence will be replaced by the finished sequence as soon as it is  
 available and the accession number will be preserved.  
 \* NOTE: This is a 'working draft' sequence.  
 \* This sequence will be replaced  
 \* by the finished sequence as soon as it is available and  
 \* the accession number will be preserved.

FEATURES  
 source

1. .117962  
 /organism="Oryza sativa (japonica cultivar-group)"  
 /cultivar="Nipponbare"  
 /db\_xref="taxon:39947"  
 /chromosome="8"  
 /clone="OJ1521\_G02"  
 33654 a 24466 c 24744 g 35098 t

BASE COUNT  
 ORIGIN

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 Best Local Similarity 95.0%; Pred. No. 1.1e+03;  
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CGCGTCGGTCGACGACCTGGA 20  
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Db 117065 CGCGACGTCGACGACCTGA 117046

RESULT 12

AP004632

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

AP004632 154255 bp DNA linear HTG 21-MAR-2002  
 Oryza sativa (japonica cultivar-group) chromosome 8 clone P0623F08,  
 \*\*\* SEQUENCING IN PROGRESS \*\*\* in ordered pieces.

AP004632 1 GI:18182012  
 HTG; HTGS\_PHASE2.  
 Oryza sativa (japonica cultivar-group) (cultivar:Nipponbare) DNA,  
 clone:P0623F08.

ORGANISM Oryza sativa (japonica cultivar-group)  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Ehrhartoideae; Oryzaceae; Oryza.

REFERENCE 1  
AUTHORS Sasaki, T., Matsumoto, T. and Yamamoto, K.  
TITLE Oryza sativa nipponbare (GA3) genomic DNA, chromosome 8, PAC  
clone: P0623F08  
JOURNAL Published Only in Database (2002)  
REFERENCE 2 (bases 1 to 154255)  
AUTHORS Sasaki, T., Matsumoto, T. and Yamamoto, K.  
TITLE Direct Submission  
JOURNAL Submitted (16-JAN-2002) Takuji Sasaki, National Institute of  
Agrobiological Sciences, Rice Genome Research Program; Kamondai  
2-1-2, Tsukuba, Ibaraki 305-8602, Japan  
(E-mail: tsasaki@affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/,  
Tel: 81-298-38-7441, Fax: 81-298-38-7468)

COMMENT  
NOTE: It currently consists of 1 contigs. Gaps between the contigs  
are represented as runs of N. The order of the pieces is believed  
to be correct as given, however the sizes of the gaps between them  
are based on estimates that have provided by the submitter. This  
sequence will be replaced by the finished sequence as soon as it is  
available and the accession number will be preserved.  
\* NOTE: This is a 'working draft' sequence.  
\* This sequence will be replaced  
\* by the finished sequence as soon as it is available and  
\* the accession number will be preserved.

FEATURES  
source  
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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CGCGTCGGTGCAGGACGTGA 20  
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Db 115307 CGCGAGGTGCAGGACGTGA 115326

RESULT 13  
A46287  
LOCUS A46287  
DEFINITION Sequence 2 from Patent W09521927.  
ACCESSION A46287  
VERSION A46287.1 GI:2300513  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 24)  
AUTHORS Ratcliffe, P.J., Firth, J.D., Harris, A.L. and Pugh, C.W.  
TITLE TARGETING GENE THERAPY  
JOURNAL Patent: WO 9521927-A 2 17-AUG-1995;  
ISIS INNOVATION (GB)  
FEATURES  
source  
Location/Qualifiers  
1..24  
/organism="unidentified"  
/db\_xref="taxon:32644"

BASE COUNT 6 a 6 c 8 g 4 t

ORIGIN

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Best Local Similarity 100.0%; Pred. No. 2.7e+03;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
|||||

Db 7 GTGCAGGACGTGACAAAT 24  
|||||

RESULT 14  
AX023673  
LOCUS AX023673  
DEFINITION Sequence 15 from Patent WO0017371.  
ACCESSION AX023673  
VERSION AX023673.1 GI:10184034  
KEYWORDS  
SOURCE Mus sp.  
ORGANISM Mus sp.  
REFERENCE 1 (bases 1 to 24)  
AUTHORS Binley, K.M. and Naylor, S.  
TITLE Polynucleotide constructs and uses thereof  
JOURNAL Patent: WO 0017371-A 15 30-MAR-2000;  
BINLEY KATIE MARY (GB); NAYLOR STUART (GB); OXFORD BIOMEDICA LTD (GB)

FEATURES  
source  
Location/Qualifiers  
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/organism="Mus sp."  
/db\_xref="taxon:10095"

BASE COUNT 6 a 6 c 8 g 4 t

ORIGIN

Query Match 72.0%; Score 18; DB 6; Length 24;  
Best Local Similarity 100.0%; Pred. No. 2.7e+03;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
|||||

Db 7 GTGCAGGACGTGACAAAT 24  
|||||

RESULT 15  
AX048713  
LOCUS AX048713  
DEFINITION Sequence 13 from Patent WO0069908.  
ACCESSION AX048713  
VERSION AX048713.1 GI:12225858  
KEYWORDS  
SOURCE Mus sp.  
ORGANISM Mus sp.  
REFERENCE 1 (bases 1 to 24)  
AUTHORS Ratcliffe, P.J., Maxwell, P.H. and Pugh, C.W.  
TITLE Interaction between the vhl tumour suppressor and hypoxia inducible  
factor, and assay methods relating thereto  
JOURNAL Patent: WO 0069908-A 13 23-NOV-2000;  
ISIS INNOVATION LIMITED (GB)  
FEATURES  
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Location/Qualifiers  
1..24  
/organism="Mus sp."  
/db\_xref="taxon:10095"

BASE COUNT 6 a 6 c 8 g 4 t

ORIGIN

Query Match 72.0%; Score 18; DB 6; Length 24;  
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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
|||||

Db 7 GTGCAGGACGTGACAAAT 24  
|||||

Search completed: January 3, 2003, 23:53:01  
Job time : 316.158 secs

GenCore version 5.1.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:02 ; Search time 41.7981 Seconds  
(without alignments)  
1346.950 Million cell updates/sec

Title: US-09-787-562-1

Perfect score: 25

Sequence: 1 cgcgcggtgcaggacgtgacaaat 25

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	25	100.0	25	21	AA111993
2	19	76.0	19	21	AA111994
3	18.6	74.4	3670	17	AAT36388
4	18.6	74.4	3670	20	AAV83129
5	18.6	74.4	3670	21	AAZ38241
6	18	72.0	24	16	AAQ94548
7	18	72.0	24	20	AAZ11422
8	18	72.0	24	21	AA12007
9	18	72.0	24	22	AAC88980
					Murine PKG HRE P42
					Murine PKG HRE tru
					Periplasmic Beta-N
					Hexosaminidase enz
					Vibrio furnissii e
					Hypoxia-inducible
					Hypoxia responsive
					Murine HRE mpkg DN
					Murine hypoxic res

10	18	72.0	513	22	AAH20729	Murine phosphoglyc
11	18	72.0	4768	22	AAH20729	Plasmid vector pbg
12	18	72.0	4768	24	AAH20729	Nucleotide sequenc
13	18	72.0	4768	24	AAH20729	Plasmid pbg2 vecto
14	18	72.0	4768	24	AAH20729	Gene targeting ve
15	18	72.0	4847	22	AAH20729	PGK-crc-pA vector
16	18	72.0	5365	22	AAH20729	Retroviral vector
17	18	72.0	5377	21	AAH20729	Expression vector
18	18	72.0	5581	22	AAH20729	Exprt gene containi
19	18	72.0	6355	22	AAH20729	Plasmid vector pbg
20	18	72.0	6355	24	AAH20729	Nucleotide sequenc
21	18	72.0	6355	24	AAH20729	Plasmid pbg4 vecto
22	18	72.0	6355	24	AAH20729	Gene targeting ve
23	18	72.0	7090	22	AAH20729	Plasmid pbg4 vecto
24	18	72.0	7617	18	AAH20729	Plasmid pbg4 vecto
25	18	72.0	8388	15	AAH20729	Plasmid pbg4 vecto
26	18	72.0	9725	21	AAH20729	Plasmid pbg4 vecto
27	18	72.0	9732	21	AAH20729	Plasmid pbg4 vecto
28	18	72.0	9738	21	AAH20729	Plasmid pbg4 vecto
29	18	72.0	9873	21	AAH20729	Plasmid pbg4 vecto
30	18	72.0	10054	21	AAH20729	Plasmid pbg4 vecto
31	18	72.0	11162	24	AAH20729	Plasmid pbg4 vecto
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33	18	72.0	11162	24	AAH20729	Plasmid pbg4 vecto
34	18	72.0	11162	24	AAH20729	Plasmid pbg4 vecto
35	18	72.0	13928	22	AAH20729	Plasmid pbg4 vecto
36	18	72.0	15692	20	AAH20729	Plasmid pbg4 vecto
37	18	72.0	15692	20	AAH20729	Plasmid pbg4 vecto
38	18	72.0	15701	20	AAH20729	Plasmid pbg4 vecto
39	17.6	70.4	4256	19	AAH20729	Plasmid pbg4 vecto
40	17.6	70.4	4256	20	AAH20729	Plasmid pbg4 vecto
41	17.4	69.6	4568	23	AAH20729	Plasmid pbg4 vecto
42	17.4	69.6	7108	23	AAH20729	Plasmid pbg4 vecto
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44	17.2	68.8	4725	21	AAH20729	Plasmid pbg4 vecto
45	17.2	68.8	4737	21	AAH20729	Plasmid pbg4 vecto

#### ALIGNMENTS

#### RESULT 1

AA111993  
ID AA111993 standard; DNA; 25 BP.

XX AA111993;

XX 14-AUG-2000 (first entry)

DT Murine PKG HRE P42 DNA sequence.

DE HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;

KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;

KW cardiovascular disease; peripheral arterial disease; cancer;

KW phosphoglycerate kinase; PGK; murine; ds.

OS Mus sp.

XX WO200017371-A1.

XX 30-MAR-2000.

XX 22-SEP-1999; 99WO-GB03181.

XX 23-SEP-1998; 98WO-GB02885.

XX 28-JAN-1999; 99GB-0001906.

XX 16-FEB-1999; 99GB-0003538.

XX (OXFO-) OXFORD BIOMEDICA UK LTD.

XX Binley KM, Naylor S;

XX WPI; 2000-283595/24.

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XX Novel polynucleotide constructs comprising at least two repeats of a
PT hypoxia response element useful for driving expression of nucleic acids
PT of interest in a cell under hypoxic conditions
XX
XX Disclosure: Page 11; 155pp; English.
XX
XX This invention describes novel polynucleotide comprising at least 2
CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible
CC factor (HIF) consensus binding sites within each of the 2 repeats are
CC separated by a spacer of at least 20 contiguous nucleotides. The products
CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic
CC activity and can be used for gene therapy. The polynucleotides are useful
CC for delivering nucleic acids of interest to mammalian cells. Lentiviral
CC vectors are responsive to hypoxic agents and to agents that mimic
CC hypoxia. This regulation can be harnessed in vitro to enhance the
CC production of the vector and can be used in vivo to regulate gene
CC expression in response to a physiological signal. The vectors have
CC utility in disease, where ischaemia, including hypoxia, is a feature,
CC e.g. cardiovascular disease, peripheral arterial disease, cancer and
CC arthritis. The novel regulatory construct is capable of driving very high
CC levels of transcription under conditions of hypoxia whilst providing only
CC low basal levels of transcription under normal oxygen conditions. The
CC polynucleotide construct targets cells within a tumor mass that are under
CC conditions of hypoxia without affecting normal surrounding tissue. This
CC polynucleotide construct targets cells within a tumor mass that are under
CC conditions of hypoxia without affecting normal surrounding tissue. This
CC sequence represents a murine phosphoglycerate kinase (PGK) HRE P24 DNA
CC fragment as described in the method of the invention.
XX
SQ Sequence 25 BP; 6 A; 6 C; 9 G; 4 T; 0 other;
Query Match 100.0%; Score 25; DB 21; Length 25;
Best Local Similarity 100.0%; Pred. No. 0.034;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CGCGTCGGTGCAGGACGTGACAAAT 25
DB 1 CGCGTCGGTGCAGGACGTGACAAAT 25
RESULT 2
AA11994
ID AA11994 standard; DNA; 19 BP.
AC AA11994;
XX
XX 14-AUG-2000 (first entry)
DE Murine PGK HRE truncated P18 DNA sequence.
XX
XX HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;
KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;
KW cardiovascular disease; peripheral arterial disease; cancer;
KW phosphoglycerate kinase; PGK; murine; ds.
XX
XX Mus sp.
XX WO200017371-A1.
XX
XX 30-MAR-2000.
XX
XX 22-SEP-1999; 99WO-GB03181.
XX
XX 23-SEP-1998; 98WO-GB02885.
XX 28-JAN-1999; 99GB-0001906.
XX 16-FEB-1999; 99GB-0003538.
XX
XX (OXFO-) OXFORD BIOMEDICA UK LTD.
XX
XX Binley KM, Naylor S;
XX WPI; 2000-283595/24.
XX
XX Novel polynucleotide constructs comprising at least two repeats of a
PT

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PT hypoxia response element useful for driving expression of nucleic acids
PT of interest in a cell under hypoxic conditions
XX
XX Disclosure: Page 11; 155pp; English.
XX
XX This invention describes novel polynucleotide comprising at least 2
CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible
CC factor (HIF) consensus binding sites within each of the 2 repeats are
CC separated by a spacer of at least 20 contiguous nucleotides. The products
CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic
CC activity and can be used for gene therapy. The polynucleotides are useful
CC for delivering nucleic acids of interest to mammalian cells. Lentiviral
CC vectors are responsive to hypoxic agents and to agents that mimic
CC hypoxia. This regulation can be harnessed in vitro to enhance the
CC production of the vector and can be used in vivo to regulate gene
CC expression in response to a physiological signal. The vectors have
CC utility in disease, where ischaemia, including hypoxia, is a feature,
CC e.g. cardiovascular disease, peripheral arterial disease, cancer and
CC arthritis. The novel regulatory construct is capable of driving very high
CC levels of transcription under conditions of hypoxia whilst providing only
CC low basal levels of transcription under normal oxygen conditions. The
CC polynucleotide construct targets cells within a tumor mass that are under
CC conditions of hypoxia without affecting normal surrounding tissue. This
CC sequence represents a murine phosphoglycerate kinase (PGK) HRE truncated
CC P18 DNA fragment as described in the method of the invention.
XX
SQ Sequence 19 BP; 4 A; 4 C; 8 G; 3 T; 0 other;
Query Match 76.0%; Score 19; DB 21; Length 19;
Best Local Similarity 100.0%; Pred. No. 20;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 4 GTCGGTGCAGGACGTGACA 22
DB 1 GTCGGTGCAGGACGTGACA 19
RESULT 3
AAT36388
ID AAT36388 standard; DNA; 3670 BP.
XX
XX AAT36388;
AC AAT36388;
XX
XX 14-JAN-1997 (first entry)
DE Periplasmic Beta-N-acetylglucosaminidase coding sequence.
XX
XX Periplasmic Beta-N-acetylglucosaminidase; periplasmic Beta-N-acetylglucosaminidase;
KW Beta-N-acetylglucosaminidase; chitin; oligosaccharide; catabolic;
KW catabolism; ss.
XX
XX Vibrio furnissii.
XX
XX Key Location/Qualifiers
FT CDS 845..2680
FT /*tag= a
FT /product= Periplasmic Beta-N-acetylglucosaminidase
XX
XX WO9625424-A1.
XX
XX 22-AUG-1996.
XX
XX 13-FEB-1996; 96WO-US02332.
XX
XX 13-FEB-1995; 95US-0386727.
XX
XX (UJJO ) UNIV JOHNS HOPKINS.
XX
XX Bassler B, Chitlaru E, Keyhani N, Roseman S, Rowe C;
XX Yu C;
XX WPI; 1996-393335/39.
XX P-PSDB; AAW02157.
XX

```

XX Chitin biosynthetic enzymes end I, exo I and exo II - are  
PT periplasmic chito:dextrinase(s), periplasmic beta-GlcNAcidase(s) and  
PT aryl beta-N-acetyl:gluco:amidase(s), respectively  
XX  
PS Claim 12; Page 71-73; 101pp; English.  
XX  
CC Periplasmic chitodextrinase (AAW02156), periplasmic  
CC Beta-N-acetylglucosaminidase (AAW02157) and aryl  
CC Beta-N-acetylglucosaminidase (AAW02158) can be used to produce chitin  
CC oligosaccharides with the structure (GlcNAc)<sub>n</sub> where n is 2 or  
CC higher, by contacting them with soluble chitin. The enzymes are  
CC encoded by the genes endI, exoI and exoII respectively. They are  
CC all genes involved in the catabolic pathway of chitin.  
XX  
SQ Sequence 3670 BP; 858 A; 1046 C; 958 G; 808 T; 0 other;  
Query Match 74.4%; Score 18.6; DB 17; Length 3670;  
Best Local Similarity 84.0%; Pred. No. 47;  
Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
XX  
QY 1 CGCGTCGGTGCGAGCGTGACAAAT 25  
Db 1033 CAGCCGGGAGCAGCGTGATAAAT 1057  
RESULT 4  
AAV83129  
ID .AAV83129 standard; cDNA; 3670 BP.  
XX  
AC AAV83129;  
XX  
XX 02-MAR-1999 (first entry)  
DT 08-MAR-1999 (revised)  
XX  
XX Hexosaminidase enzyme coding sequence.  
DE  
XX  
XX Hexosaminidase; enzyme; laundry; cleaning agent; hydrolysis;  
KW anti-microbial; detergent; surfactant; ss.  
XX  
OS Unidentified.  
XX  
XX WO980512-A1.  
PN  
XX  
PD 12-NOV-1998.  
XX  
XX 05-MAY-1998; 98WO-US09125.  
PF  
XX 19-AUG-1997; 97US-0056132.  
PR 06-MAY-1997; 97US-0045756.  
XX  
XX (PROC ) PROCTER & GAMBLE CO.  
PA  
XX  
PI Convents AC, Moese RL, Wolff AM;  
XX  
XX WPI; 1999-024116/02.  
DR P-PSDB; AAW85599.  
XX  
XX Laundry and cleaning compositions containing hexosaminidase - to  
PT provide antimicrobial activity and remove biofilm  
XX  
XX Claim 2; Page 42-44; 64pp; English.  
XX  
XX Novel hexosaminidase enzymes (AAW85599-605) can be used in  
CC combination in an aqueous laundry or cleaning product. The cleaning  
CC product is used especially used to launder fabrics and to clean  
CC dishes and tableware, particularly in an automatic dishwasher, but  
CC may also be used generally as hard surface cleaner. The cleaning  
CC product imparts antimicrobial activity and/or eliminates biofilm,  
CC the hexosaminidases having a minimum inhibitory concentration of  
CC less than about 0.125% but more preferably less than about 0.025%.  
CC (NB: entry was revised to change incorrect cross references in  
CC Comments field).

XX  
SQ Sequence 3670 BP; 858 A; 1046 C; 958 G; 808 T; 0 other;  
Query Match 74.4%; Score 18.6; DB 20; Length 3670;  
Best Local Similarity 84.0%; Pred. No. 47;  
Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
XX  
QY 1 CGCGTCGGTGCGAGCGTGACAAAT 25  
Db 1033 CAGCCGGGAGCAGCGTGATAAAT 1057  
RESULT 5  
AAZ38241  
ID .AAZ38241 standard; DNA; 3670 BP.  
XX  
AC AAZ38241;  
XX  
XX 09-FEB-2000 (first entry)  
DT  
XX  
XX Vibrio furnissii exoI gene, encoding periplasmic beta-GlcNAcidase.  
DE  
XX  
XX Beta-GlcNAcidase; periplasmic; exoI; Exo-I; chitin; chitosan; exoenzyme;  
KW beta 1-4 N-acetylglucosamine; GlcNAc; degradation; catabolism;  
KW oligosaccharide; agriculture; medicine; nitrogen fixing nodules;  
KW disease resistance; induction; fungicide; antimetastatic;  
KW Lewis lung carcinoma; immune system; macrophage; activation; production;  
KW recombinant protein; ss.  
XX  
XX Vibrio furnissii.  
OS  
XX  
XX  
FH Key Location/Qualifiers  
FT CDS 845..2680  
FT /\*tag= a  
FT /product= "Vibrio furnissii periplasmic beta-GlcNAcidase  
FT (Exo-I)"  
XX  
XX US5985644-A.  
XX  
XX 16-NOV-1999.  
XX  
XX 13-FEB-1996; 96US-0600452.  
XX  
XX 13-FEB-1995; 95US-0386727.  
XX  
XX (UYJO ) UNIV JOHNS HOPKINS.  
XX  
XX Bassler B, Chitlaru E, Yu C, Roseman S, Keyhani NO;  
XX WPI; 2000-022280/02.  
XX P-PSDB; AAY52305.  
XX  
XX DNA encoding periplasmic chitodextrinase endoenzyme -  
XX  
XX Example 2; Columns 39-42; 37pp; English.  
XX  
XX This sequence represents Vibrio furnissii exoI gene, which encodes  
CC periplasmic beta-GlcNAcidase (Exo-I). Chitin is a homopolymer of beta  
CC 1-4 N-acetylglucosamine (GlcNAc). Chitin degradation by V. furnissii  
CC involves several signal transducing systems and a multitude of proteins,  
CC in contrast to other organisms which only require two enzymes to degrade  
CC chitin to GlcNAc. Exo-I is an exoenzyme, cleaving off terminal GlcNAc  
CC units from higher chitin-derived oligosaccharides to produce GlcNAc  
CC monomers and dimers. Chitin oligosaccharides have been recently shown  
CC to be physiologically active and are useful in agriculture and medicine.  
CC Derivatized oligosaccharides are generated by Rhizobium species as  
CC signals for the formation of nitrogen fixing nodules by leguminous plants  
CC and also induce disease resistance in certain plants. They also inhibit  
CC the growth of several fungal pathogens. The GlcNAc hexamer is a potent  
CC antimetastatic agent against Lewis lung carcinoma, and GlcNAc polymers of  
CC varying lengths activate macrophages and the immune system. Prior art  
CC methods of production of such oligosaccharides are prohibitively  
CC expensive and there are limitations in resolving mixtures of these

CC compounds. Use of recombinant V. furnissii chitin catabolic enzymes  
CC may help to reduce or eliminate these problems.

XX Sequence 3670 BP; 858 A; 1046 C; 958 G; 808 T; 0 other;

XX Query Match 74.4%; Score 18.6; DB 21; Length 3670;  
XX Best Local Similarity 84.0%; Pred. No. 47;  
XX Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CCGCTCGTGCAGGACGTGACAAAT 25  
DB 1033 CACGCCGAGCAGGACGTGATAAAT 1057

RESULT 6  
AAQ99458  
ID AAQ99458 standard; DNA; 24 BP.

XX AC AAQ99458;

DT 19-MAR-1996 (first entry)

XX Hypoxia-inducible phosphoglycerate kinase-1 expression control sequence.

XX Hypoxia; response element; erythropoietin; phosphoglycerate kinase;  
KW PGK-1; gene therapy; tumour; cancer; P18; P24; ss.

XX Mus sp.

XX W09521927-A2.

XX 17-AUG-1995.

XX 15-FEB-1995; 95WO-GB003322.

XX 15-FEB-1994; 94GB-0002857.

XX (ISIS-) ISIS INNOVATION LTD.

PI Firth JD, Harris AL, Pugh CW, Ratcliffe PJ;

DR WPI; 1995-293128/38.

XX Novel method of targeting gene therapy - using a hypoxically  
PT inducible expression control sequence linked to a species active  
PT against disease

XX Disclosure; Page 4; 30pp; English.

XX AAQ99458 and AAQ99459 are hypoxia inducible transcription control  
CC elements P24 and P18, respectively. P24 and P18 may be linked to at  
CC least one gene encoding a protein that has activity against disease e.g.  
CC CD2, CD4, etc. Such a construct may be used to treat a patient  
CC suffering from a disease in which hypoxia is a cause or symptom.  
CC The constructs are particularly used for treating tumours.

XX Sequence 24 BP; 6 A; 6 C; 8 G; 4 T; 0 other;

QY Query Match 72.0%; Score 18; DB 16; Length 24;  
DB Best Local Similarity 100.0%; Pred. No. 59;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
DB 7 GTGCAGGACGTGACAAAT 24

RESULT 7  
AAZ11422  
ID AAZ11422 standard; DNA; 24 BP.

XX AC AAZ11422;

XX

DT 26-OCT-1999 (first entry)  
XX Hypoxia responsive sequence mpck.

XX Retroviral vector; functional splice donor site; hybrid viral vector;  
KW functional splice acceptor site; in vivo gene delivery; therapeutic;  
KW lentiviral vector; modified hematopoietic stem cell; MHSC; tumour;  
KW ischemia; hypoxia response element; HRE; hypoxia; ss.

XX Mus sp.

XX W09915684-A2.

XX 01-APR-1999.

XX 23-SEP-1998; 98WO-GB02885.

XX 25-SEP-1997; 97GB-0020465.

XX 23-SEP-1997; 97GB-0020216.

XX (OXFO-) OXFORD BIOMEDICA UK LTD.

XX Bebbington C, Binley KM, Lewis C, Naylor S;

XX WPI; 1999-263482/22.

XX New retroviral vectors, for, e.g. delivering nucleotide sequences to  
PT solid tumor sites

XX Example 1 (page 68); Fig 1 (page 1/43); 288pp; English.

XX The invention relates to a retroviral vector (RVV) comprising a  
CC functional splice donor site (FSDS) and a functional splice acceptor  
CC site (FSAS) where: (i) the FSDS and the FSAS flank a first nucleotide  
CC sequence of interest (NOI); (ii) the FSDS is upstream of the FSAS; (iii)  
CC the RVV is derived from a retroviral pro-vector; (iv) the retroviral  
CC pro-vector comprises a first nucleotide sequence (NS) capable of yielding  
CC the FSDS and a second NS capable of yielding the FSAS; and (v) the first  
CC NS is downstream of the second NS, such that the RVV is formed as a  
CC result of reverse transcription of the retroviral pro-vector. A hybrid  
CC viral vector (VV) system for in vivo gene delivery, which system  
CC comprises a primary VV which encodes a secondary VV, the primary vector  
CC capable of infecting a first target cell and of expressing the secondary  
CC VV, which secondary vector is capable of transducing a secondary target  
CC cell, where the primary vector is obtainable from or is based on a  
CC adenoviral vector and the secondary VV is obtainable from or is based on  
CC a RVV preferably a lentiviral vector (LVV) is also provided. The systems  
CC can be used for delivering NOIs to one or more target sites. The NOIs may  
CC encode therapeutic or diagnostic agents. The methods are used  
CC particularly for producing modified hematopoietic stem cells (MHSCs) to  
CC deliver NOIs to sites such as solid tumours which are characterised by  
CC ischemia, such as hypoxia or low glucose concentration. The system  
CC permits the stable expression of NOIs in targeted cells, e.g. rapidly  
CC dividing cells. Sequences AAZ11420-430 represent nucleotide sequences  
CC that are responsive to hypoxia.

XX Sequence 24 BP; 6 A; 6 C; 8 G; 4 T; 0 other;

QY Query Match 72.0%; Score 18; DB 20; Length 24;  
DB Best Local Similarity 100.0%; Pred. No. 59;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
DB 7 GTGCAGGACGTGACAAAT 24

RESULT 8  
AAAL2007  
ID AAAL2007 standard; DNA; 24 BP.

XX AC AAAL2007;

XX





XX WPI; 2001-343817/36.

DR New permanent amniocyte cell lines, useful for producing viral gene

PT therapy vectors or mutant adenoviruses, express the adenoviral E1A and

PT E1B gene products -

XX

XX Example 1; Page 61; 72pp; German.

PS

XX This invention describes novel permanent amniocyte cell lines (A),

CC containing at least one nucleic acid (I) that causes expression of the

CC gene products (II) of the adenoviral E1A and E1B regions. (A) are used to

CC produce gene therapy vectors, especially adeno. adeno-associated, retro

CC or lentiviral vectors, particularly first- or second generation,

CC large-capacity or deleted adenoviral vectors. (A) are also used to

CC produce adenoviral mutants, optionally with modified tropism. The vectors

CC may express a wide range of therapeutic proteins or antisense RNAs.

CC Adenoviral mutants, unable to express the E1B 55 kDa protein, are useful

CC for treating tumors, they replicate in the cells but not significantly in

CC normal primary cells. (A) can be made efficiently, simply and

CC reproducibly. The products of the invention have cytostatic activity.

CC This sequence represents the murine phosphoglycerate kinase promoter

CC found in plasmid STK146.

XX

XX Sequence 513 BP; 79 A; 171 C; 161 G; 102 T; 0 other;

SQ

Query Match 72.0%; Score 18; DB 22; Length 513;

Best Local Similarity 100.0%; Pred. No. 76;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25

DB 221 GTGCAGGACGTGACAAAT 238

RESULT 11

AAS05243

ID AAS05243 standard; DNA; 4768 BP.

XX

XX AAS05243;

XX

XX 07-SEP-2001 (first entry)

XX

XX Plasmid vector pDG2 used as a construct for TRP genes.

XX

XX Trinucleotide repeat protein; TRP; T243; embryonic stem cell; ES; pDG2;

KW transgenic animal; knockout mouse; triplet repeat expansion;

KW fragile X syndrome; Huntington's disease; cyclic; circular; ds.

XX

OS Synthetic.

XX

XX W0200130798-Al.

XX

XX 03-MAY-2001.

XX

XX 26-OCT-2000; 2000WO-US29382.

XX

XX 26-OCT-1999; 99US-0161488.

XX

XX (DELTA-) DELTAGEN INC.

PA Klein R, Matthews W, Moore M, Allen KD;

PI WPI; 2001-300473/31.

DR

XX Novel transgenic animals useful as animal model for characterization of

PT function of a gene encoding trinucleotide repeat proteins (TRPs),

PT contains heterozygous disruption in a gene encoding TRP -

XX

PS Disclosure; Fig 2B; 106pp; English.

XX

XX The present sequence for plasmid vector pDG2 is used as a construct

CC for genes encoding trinucleotide repeat proteins (TRP) such as gene

CC

CC T243 to produce disruption in the DNA. The invention describes

CC methods of producing embryonic stem (ES) cells comprising a heterozygous

CC disruption in a target DNA sequence (preferably T243) encoding a TRP and

CC of producing a knockout mouse comprising a homozygous disruption in a

CC gene encoding TRP, where the disruption inhibits the production of the

CC wild type TRP. The invention also relates to identifying agents capable

CC of affecting a phenotype of a knockout mouse. Also described are methods

CC of determining whether expansion of the trinucleotide repeat in a gene

CC encoding TRP produces a phenotypic change. The transgenic animals and

CC the cells are useful for identifying compounds capable of ameliorating

CC disease symptoms, and as test substrates for the identification of drugs,

CC pharmaceuticals, therapies and interventions which may be effective in

CC treating trinucleotide repeat disorders e.g. fragile X syndrome and

CC Huntington's disease. The animal models for trinucleotide repeat

CC disorders are ideal model systems to study the progression of disease in

CC vivo, the molecular basis of these diseases and show the features

CC observed in human disease. Using the mice, it is possible to model both

CC the pathogenic mechanism and the trinucleotide repeat instability in the

CC mouse.

XX

SQ Sequence 4768 BP; 1124 A; 1218 C; 1269 G; 1157 T; 0 other;

Query Match 72.0%; Score 18; DB 22; Length 4768;

Best Local Similarity 100.0%; Pred. No. 92;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25

DB 2824 GTGCAGGACGTGACAAAT 2841

RESULT 12

ABL42019

ID ABL42019 standard; DNA; 4768 BP.

XX

XX ABL42019;

XX

XX 11-JUN-2002 (first entry)

XX

XX Nucleotide sequence of vector pDG2.

XX

XX pDG2; transgenic animal; matrix metalloproteinase-23 gene; MMP-23 gene;

KW ss.

XX

OS Synthetic.

XX

XX US2002023275-Al.

XX

XX 21-FEB-2002.

XX

XX 17-MAY-2001; 2001US-0861077.

XX

XX 17-MAY-2000; 2000US-204972P.

PR 29-JUN-2000; 2000US-215394P.

XX

XX (LEVI/) LEVITEN M W.

PA Leviten MW;

PI WPI; 2002-255684/30.

DR

XX Non-human transgenic animal useful as a model for disease and for

PT identifying agents that modulate gene expression and gene function,

PT comprises a disruption in the matrix metalloproteinase-23 gene -

XX

XX Example 1; Fig 2B; 38pp; English.

PS

XX The present sequence represents vector pDG2. This vector contains an

CC ampicillin resistance gene and a neomycin gene. The vector is used in

CC the invention. The specification describes a non-human transgenic animal

CC comprising a disruption in the matrix metalloproteinase (MMP)-23 gene.

CC Transgenic animals of the invention comprising a homozygous or

CC heterozygous disruption in MMP23 gene are useful for identifying agents

CC

CC which modulate MMP23 expression or function. They are also useful for  
 CC identifying agents that are capable of ameliorating a phenotype of a  
 CC transgenic animal comprising a disruption in an MMP-23 gene or  
 CC ameliorating a disease associated with the phenotype of a transgenic  
 CC animal comprising a disruption in the MMP-23 gene. The animals are  
 CC useful as an animal model for diseases, disorders and conditions  
 CC characterized by a disruption in a gene encoding a metalloproteinase,  
 CC more particularly disease, disorders and conditions associated with the  
 CC phenotypes demonstrated by the knockout mice. The transgenic animals  
 CC are useful as test substrates for identification of drugs,  
 CC pharmaceuticals and therapies effective in treating diseases, disorders  
 CC and conditions associated with disruption in the target gene. The  
 CC animal is useful for testing and developing new treatments relating  
 CC to behavioural phenotypes demonstrated by the animal models.

XX SQ Sequence 4768 BP; 1124 A; 1218 C; 1269 G; 1157 T; 0 other;  
 Query Match 72.0%; Score 18; DB 24; Length 4768;  
 Best Local Similarity 100.0%; Pred. No. 92;  
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 8 GTGCAGGACGTGACAAAT 25  
 |||||  
 Db 2824 GTGCAGGACGTGACAAAT 2841

RESULT 13  
 AAD28659  
 ID AAD28659 standard; DNA; 4768 BP.  
 AC AAD28659;  
 XX 07-MAY-2002 (first entry)  
 DE Plasmid pDG2 vector.  
 XX Plasmid pDG2; DNA construct; embryonic stem cell; cell disruption; Neo';  
 KW neomycin; ampicillin resistance gene; ds.  
 XX Unidentified.  
 OS WO200204621-A2.  
 PN 17-JAN-2002.  
 PD 11-JUL-2000; 2000WO-US18812.  
 PF 11-JUL-2000; 2000WO-US18812.  
 PR 11-JUL-2000; 2000WO-US18812.  
 PA (DELT-) DELTAGEN INC.  
 XX Klein RD, Brennan TJ;  
 PI WPI; 2002-164642/21.  
 DR Novel nucleotide construct for generating DNA constructs for  
 PT introducing into embryonic stem cell, comprising a sequence encoding a  
 PT positive selection marker flanked by restriction enzyme sites -  
 XX Claim 10; Fig 2B; 64pp; English.

CC The invention relates to nucleotide construct for generating DNA  
 CC constructs. The nucleotide construct comprises a sequence encoding a  
 CC positive selection marker flanked by restriction enzyme sites, where  
 CC restriction site is flanked by sequences which are not complementary to  
 CC each other and which do not include at least one type of base at any  
 CC position, where the construct can be treated so that single-stranded  
 CC regions are created at each sequence lacking at least one nucleotide.  
 CC The nucleotide construct is useful in a rapid and efficient method for  
 CC generating DNA constructs suitable for introduction into embryonic stem  
 CC cells and for disrupting the function of a gene in a cell. The present  
 CC sequence is plasmid pDG2 vector construct containing an ampicillin  
 CC resistance gene and neomycin gene (Neo'). On each site of the Neo' gene

CC are two sites for ligation independent cloning along with restriction  
 CC sites.

XX SQ Sequence 4768 BP; 1124 A; 1218 C; 1269 G; 1157 T; 0 other;  
 Query Match 72.0%; Score 18; DB 24; Length 4768;  
 Best Local Similarity 100.0%; Pred. No. 92;  
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
 |||||  
 Db 2824 GTGCAGGACGTGACAAAT 2841

RESULT 14  
 AAS17143  
 ID AAS17143 standard; DNA; 4768 BP.  
 XX AAS17143;  
 AC AAS17143;  
 XX 14-FEB-2002 (first entry)  
 DE Gene targeting vector pDG2.  
 XX

KW pDG2; ds; retina-specific nuclear receptor; gene targeting;  
 KW lymphoid-specific GPCR; melanocyte stimulating hormone receptor;  
 KW magnesium-dependent protein phosphatase; transgenic animal;  
 KW chemokine receptor 1-like protein; CGMP phosphodiesterase;  
 KW sulfotransferase gene; tumour; cancer; retinal degeneration;  
 KW retinitis pigmentosa.

XX Escherichia coli.  
 OS Synthetic.  
 OS WO200167855-A2.

XX 20-SEP-2001.  
 XX 16-MAR-2001; 2001WO-US08664.  
 XX 16-MAR-2000; 2000US-190348P.  
 PR 22-MAR-2000; 2000US-191128P.  
 PR 22-MAR-2000; 2000US-191129P.  
 PR 22-MAR-2000; 2000US-191142P.  
 PR 22-MAR-2000; 2000US-191235P.  
 PR 22-MAR-2000; 2000US-191336P.  
 PR 22-MAR-2000; 2000US-191400P.  
 PR 15-MAY-2000; 2000US-204227P.  
 PR 15-MAY-2000; 2000US-204230P.  
 PR 29-JUN-2000; 2000US-215214P.  
 PR 06-JUL-2000; 2000US-216249P.  
 PR 06-JUL-2000; 2000US-216264P.  
 PR 06-JUL-2000; 2000US-216765P.  
 PR 12-JUL-2000; 2000US-218075P.  
 PR 19-JUL-2000; 2000US-219167P.  
 PR 19-JUL-2000; 2000US-219182P.  
 PR 27-JUL-2000; 2000US-221485P.  
 PR 07-AUG-2000; 2000US-223173P.

XX (DELT-) DELTAGEN INC.

XX Allen KB, Guenther C, Phillips R;

XX WPI; 2002-041167/05.

XX New targeting construct comprising a first and a second polynucleotide  
 PT homologous to a target gene, and a selectable marker, useful for  
 PT introducing targeted mutations into embryonic cells -

XX Example 3; Fig 2B; 105pp; English.

CC The invention relates to a targeting construct comprising two sequences  
 CC homologous to a target gene, and a selectable marker, is new. The target

CC gene is a retina-specific nuclear receptor gene, a lymphoid-specific GPCR  
CC (G protein coupled receptor) gene, a melanocyte stimulating hormone  
CC receptor gene, a magnesium-dependent protein phosphatase gene, chemokine  
CC receptor 1-like protein gene, a cGMP phosphodiesterase gene, or a  
CC sulfolipase gene. Also included are transgenic mice comprising a  
CC disruption in a target gene, where the mouse exhibits an eye abnormality,  
CC cellular infiltration, hypoaffective behaviour, lung abnormality, elevated  
CC white blood cell count, abnormality in the aorta, kidney, liver, lymph  
CC nodes, skin or salivary gland, increased body and organ weight, or  
CC elevated levels of ALT (not defined), phosphorus, potassium, or  
CC bilirubin, aggressive, hyperactive, increased activity or decreased  
CC anxiety behaviour. The construct is used for introducing targeted  
CC mutations into embryonic cells. The animal and cell-based systems may be  
CC used as models for diseases or conditions associated with physiological,  
CC histological or behavioural phenotypes relating to a disruption in a  
CC target gene (e.g. tumours, cancer, retinal degeneration and retinitis  
CC pigmentosa) and in screening or identifying compounds capable of  
CC ameliorating or treating diseases. The present sequence is the vector  
CC pDG2 used to generate the gene targeting construct of the invention.  
XX  
SQ Sequence 4768 BP; 1124 A; 1218 C; 1269 G; 1157 T; 0 other;

Query Match 72.0%; Score 18; DB 24; Length 4768;  
Best Local Similarity 100.0%; Pred. No. 92;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25

|||||

Db 2824 GTGCAGGACGTGACAAAT 2841

#### RESULT 15

AAD09280

ID AAD09280 standard; DNA; 4847 BP.

XX

AC AAD09280;

DT 12-SEP-2001 (first entry)

XX  
DE PGK-cre-pA vector DNA.

XX DNA recombinase domain; protein transduction domain; PTD;  
KW gene alteration; fusion protein; Human immunodeficiency virus;  
KW HIV; pgk-cre-pA vector; ds.

XX Unidentified.

XX  
PN WO200149832-A2.

XX

PD 12-JUL-2001.

XX

PF 05-JAN-2001; 2001WO-EP00060.

XX

PR 07-JAN-2000; 2000EP-0100351.

XX

PR 10-NOV-2000; 2000EP-0124595.

XX

PA (ARTE-) ARTEMIS PHARM GMBH.

XX

PI Schwenk F;

XX

XX WPI; 2001-441873/47.

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CC present invention also provides a method for inducing gene  
CC alterations in living organisms using the fusion proteins of the  
CC invention. The present sequence is a pgk-cre-pA DNA.

XX  
SQ Sequence 4847 BP; 1139 A; 1238 C; 1283 G; 1184 T; 3 other;

Query Match 72.0%; Score 18; DB 22; Length 4847;  
Best Local Similarity 100.0%; Pred. No. 92;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25

|||||

Db 2466 GTGCAGGACGTGACAAAT 2483

Search completed: January 3, 2003, 23:03:43  
Job time : 44.7981 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:17 ; Search time 332.413 Seconds  
(without alignments)  
1218.024 Million cell updates/sec

Title: US-09-787-562-1  
Perfect score: 25  
Sequence: 1 cgcgcgtgcagcagtgacaaat 25

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:\*

- 1: em\_estba:\*
- 2: em\_esthum:\*
- 3: em\_estinu:\*
- 4: em\_estinu:\*
- 5: em\_estov:\*
- 6: em\_estpl:\*
- 7: em\_estro:\*
- 8: em\_hic:\*
- 9: gb\_est1:\*
- 10: gb\_est2:\*
- 11: gb\_hic:\*
- 12: gb\_est3:\*
- 13: gb\_est4:\*
- 14: gb\_est5:\*
- 15: em\_estfun:\*
- 16: em\_estom:\*
- 17: gb\_gss:\*
- 18: em\_gss\_hum:\*
- 19: em\_gss\_inv:\*
- 20: em\_gss\_pln:\*
- 21: em\_gss\_vrt:\*
- 22: em\_gss\_fun:\*
- 23: em\_gss\_mam:\*
- 24: em\_gss\_mus:\*
- 25: em\_gss\_other:\*
- 26: em\_gss\_pro:\*
- 27: em\_gss\_rod:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES					
Result No.	Score	Query Match	Length	ID	Description
1	19.8	79.2	610	12	BG220636
2	19.2	76.8	824	12	BF253450
C 3	18.6	74.4	541	17	TA249A12Q
C 4	18.6	74.4	786	12	BG330319
C 5	17.8	71.2	516	17	AQ235963
C 6	17.8	71.2	865	17	CNS04REF
					AL294720 Tetraodon

C 7	17.8	71.2	1031	17	CNS04F4W
8	17.6	70.4	199	10	BE190622
C 9	17.6	70.4	412	17	AQ647334
C 10	17.6	70.4	414	17	AQ641517
C 11	17.6	70.4	442	17	AZ247056
C 12	17.6	70.4	449	10	BB839535
C 13	17.6	70.4	464	17	AQ844401
C 14	17.6	70.4	515	9	AA957358
C 15	17.6	70.4	564	10	AW664262
C 16	17.6	70.4	577	13	BI443703
C 17	17.6	70.4	578	9	AA386752
C 18	17.6	70.4	592	9	AI981217
C 19	17.6	70.4	615	9	AJ393812
C 20	17.6	70.4	636	17	AG063686
C 21	17.6	70.4	648	10	AV609456
C 22	17.6	70.4	673	14	BQ608828
C 23	17.6	70.4	688	17	BH670819
C 24	17.6	70.4	690	9	AJ447187
C 25	17.6	70.4	714	9	AJ449984
C 26	17.6	70.4	735	9	AJ447188
C 27	17.6	70.4	740	17	BH446078
C 28	17.6	70.4	765	9	AJ456336
C 29	17.6	70.4	766	9	AJ452191
C 30	17.6	70.4	816	10	AW983486
C 31	17.6	70.4	851	17	BH716762
C 32	17.6	70.4	972	12	BG461126
C 33	17.4	69.6	584	10	AW126461
C 34	17.4	69.6	741	13	BM073359
C 35	17.4	69.6	928	13	BI452891
C 36	17.4	69.6	1021	17	CNS05218
C 37	17.4	69.6	1722	11	AY103550
C 38	17.2	68.8	220	13	BJ266315
C 39	17.2	68.8	289	9	AI396788
C 40	17.2	68.8	393	9	AI331186
C 41	17.2	68.8	437	9	AJ432011
C 42	17.2	68.8	448	10	BB839526
C 43	17.2	68.8	455	9	AI332263
C 44	17.2	68.8	479	9	AA547770
C 45	17.2	68.8	480	9	AJ478558

## ALIGNMENTS

RESULT 1	BG220636	610 bp	mRNA	linear	EST 21-APR-2001
LOCUS	RST40423	Athersys	RAGE Library	Homo sapiens	CDNA, mRNA sequence.
DEFINITION	BG220636				
ACCESSION	BG220636				
VERSION	BG220636.1	GI:13746657			
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
AUTHORS	Harrington, J.J., Sherif, B., Rundlett, S., Jackson, P.D., Perry, R., Lerner, L., Costanzo, D., McElligott, K., Booser, S., Mays, R., Smith, E., Veloso, N., Klika, A., Hess, J., Cothren, K., Lo, K., Offenbacher, J., Danzig, J., and Ducar, M.				
TITLE	Creation of genome-wide protein expression libraries using random activation of gene expression				
JOURNAL	Nat. Biotechnol. 19 (5), 440-445 (2001)				
MEDLINE	21227151				
COMMENT	Contact: Scott J. Cain Athersys, Inc. 3201 Carnegie Ave, Cleveland, OH 44115, USA Tel: 216 431 9900 Fax: 216 361 9596 Email: scain@atersys.com High quality sequence stop: 372. Location/Qualifiers				

AL287897 Tetraodon  
BE190622 so20d12.y  
AQ647334 RPC193-EC  
AQ641517 RPC193-EC  
AZ247056 RPC1-23-4  
BB839535 BB839535  
AQ844401 an36h09.J  
AA957358 UI-R-EI-f  
AW664262 hi08e10.x  
BI443703 dai93f02.  
AA386752 vcl19q08.r  
AI981217 pat.pk004  
AJ393812 AJ393812  
AG063686 Pan trogl  
AV609456 AV609456  
BQ608828 BRY\_4742  
BH670819 BOMAT29TF  
AJ447187 AJ447187  
AJ449984 AJ449984  
AJ447188 AJ447188  
BH446078 BQFG94TR  
AJ456336 AJ456336  
AJ452191 AJ452191  
AW983486 HVSMEQ001  
BH716762 BOMBI11TF  
BG461126 RST43862  
AW126461 614072E05  
BM073359 MEST65-C1  
BI452891 603170232  
AI318185 Tetraodon  
AY103550 Zee mays  
BJ266315 BJ266315  
AI396788 fb14b03.y  
AI331186 fb05h08.y  
AJ432011 AJ432011  
BB839526 BB839526  
AI332263 fa97e01.y  
AA547770 EST188679  
AJ478558 AJ478558

Mon Jan 6 15:20:17 2003

```

source
1. .610
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Athensys RAGE Library"
/cell_line="HT1080"
/notes="See 'Creation of Genome-wide Protein Expression
Libraries using Random Activation of Gene Expression',
Nature Biotechnology, in press. Note that even though the
cell type indicated is HT1080, since a random activation
method was used, these sequence tags are not necessarily
expressed in HT1080 under normal circumstances."
BASE COUNT 113 a 206 c 186 g 104 t 1 others
ORIGIN

Query Match 79.2%; Score 19.8; DB 12; Length 610;
Best Local Similarity 91.3%; Pred. No. 3.3e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Caps 0;

QY 1 CGCGTCGGTGCAGGACGTGACAA 23
||||| ||||||| |||||||
Db 321 CGCGTCGGTGCAGGACGTGACAA 343

RESULT 2
BF253450 824 bp mRNA linear EST 22-OCT-2001
LOCUS HVSMEF0001H07f Hordeum vulgare seedling root EST library HVCDNA0007
DEFINITION (Etisolated and unstressed) Hordeum vulgare cDNA clone
HVSMEF0001H07f, mRNA sequence.
ACCESSION BF253450
VERSION BF253450.2 GI:13116515
KEYWORDS EST.
SOURCE Hordeum vulgare.
ORGANISM Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae
; Triticeae; Hordeum.
1 (bases 1 to 824)
Wing, R., Close, T.J., Klein, R., Rambo, T., Simmons, J., Choi, D.W., Fenton
Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
R.D., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex unstressed seedling root cDNA library
Unpublished (2001)
On Nov 16, 2000 this sequence version replaced gi:11182651.
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hg bases = 118
Seq primer: AATAACCCCTCACTAAAGGG
High quality sequence start: 10
High quality sequence stop: 183.
Location/Qualifiers
1. .824
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEF0001H07f"
/cdona_lib="Hordeum vulgare seedling root EST library
HVCDNA0007 (Etisolated and unstressed)"
/tissue_type="Seedling root"
/lab_host="TJCI21"
/notes="Vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and cefotaxime in covered
crystallization dishes. Five-day old seedling roots were
then harvested, total RNA was prepared, poly(A) RNA was
purified, one primary unamplified cDNA library was made,

```

and 1 million pfu were in vivo excised to give pBluescript SK(-) cDNA phagemids. These steps were performed in the TJ Close laboratory at the University of California, Riverside (Choi, Close, Fenton). Phagemids were plated and picked at the Clemson University Genomics Institute (CUGI) (Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA preparations, DNA sequencing and sequence analysis were performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main). The sequence has been trimmed to remove vector sequence and contains a minimum of 100 bases of phred value 20 or above. For more details on library preparation and sequence analysis see <http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> see Close TJ, Wing R, Klein, R., Rambo, T., Simmons, J., Choi, D.W., Fenton Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton R.D., Oates, R. and Main, D. Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html>)

```

BASE COUNT 163 a 249 c 254 g 158 t
ORIGIN

```

```

Query Match 76.8%; Score 19.2; DB 12; Length 824;
Best Local Similarity 87.5%; Pred. No. 6.1e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Caps 0;

```

```

QY 1 CGCGTCGGTGCAGGACGTGACAA 24
||||| ||||||| |||||||
Db 201 CGCGTCGGTGCAGGACGTGACAA 224

```

```

RESULT 3
TA249A120/c 541 bp DNA linear GSS 13-DEC-2000
LOCUS T. brucei sheared genomic DNA clone 249a12, reverse sequence,
DEFINITION genomic survey sequence.
ACCESSION AL482071
VERSION AL482071.1 GI:11848076
KEYWORDS GSS.
SOURCE Trypanosoma brucei.
ORGANISM Trypanosoma brucei
Eukaryota; Euzlenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
1 (bases 1 to 541)
Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
Melville, S.E., Rajandream, M.A. and Barrell, B.G.
Direct Submission
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
nhs@sanger.ac.uk
Constructed at the Institute for Genomic Research (TIGR),
Rockville, MD. Genomic DNA isolated from a cloned population of
Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared
to give a tight size distribution (
4 kb). The v + i method used for the library construction is
described in detail in Smith, H. and Venter, J.C. (Making small
insert libraries for whole genome shotgun sequencing projects. In
Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
Barrell, Oxford University Press, 1999).
Email: nhs@sanger.ac.uk
Details of T. brucei sequencing at the Sanger Centre are available
at http://www.sanger.ac.uk/projects/T\_brucei/.
Location/Qualifiers
1. .541
/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="249a12"

```

```

FEATURES
source

```

```

BASE COUNT 101 a 154 c 142 g 144 t
ORIGIN

```

```

Query Match 74.4%; Score 18.6; DB 17; Length 541;

```

Best Local Similarity 84.0%; Pred. No. 1e+03; Mismatches 0; Gaps 0;  
Matches 21; Conservative 0;

QY 1 CGCGTCGGTGCAGGCGTCACAAAT 25  
Db 330 CGCGCGGTGCAGGATCTGAAAAAT 306

## RESULT 4

BG330319/c  
LOCUS 786 bp mRNA linear EST 27-FEB-2001  
DEFINITION 602430347f1 NIH\_MGC\_18 Homo sapiens cDNA clone IMAGE:4548275 5',  
mRNA sequence.

ACCESSION BG330319  
VERSION BG330319.1 GI:13136757  
KEYWORDS EST.  
SOURCE human.

## ORGANISM

Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

## REFERENCE

AUTHORS NIH-MGC http://mgc.nci.nih.gov/.  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgapbs-femail.nih.gov

Tissue Procurement: DCTD/DTF/Gazdar  
cDNA Library Preparation: Ling Hong/Rubin Laboratory

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov  
Plate: LCM1237 row: m column: 12

High quality sequence stop: 544.  
Location/Qualifiers  
1..786

## FEATURES

source

/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:4548275"  
/clone\_lib="NIH\_MGC\_18"  
/lab\_host="DH10B (phage-resistant)"  
/tissue\_type="large cell carcinoma"  
/note="Organ: lung; Vector: pOTB7; Site1: XhoI; Site2:  
ECORI; cDNA made by oligo-dt priming. Directionally cloned  
into ECORI/XhoI sites using the following 5' adaptor:  
GCACGAG(G). Library constructed by Ling Hong in the  
laboratory of Gerald M. Rubin (University of California,  
Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and  
Superscript II RT (Life Technologies). Note: this is a  
NIH\_MGC Library."

BASE COUNT 138 a 266 c 247 g 135 t  
ORIGIN

## Query Match

Best Local Similarity 74.4%; Score 18.6; DB 12; Length 786;

Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CGCGTCGGTGCAGGCGTCACAAAT 25  
Db 700 CGCGCGGTGCAGGCGTGCACAT 676

## RESULT 5

AQ235963/c  
LOCUS 516 bp DNA linear GSS 29-SEP-1998  
DEFINITION HS\_2056\_AL\_D06\_27 CIT Approved Human Genomic Sperm Library D Homo  
sapiens genomic clone Plate=2056 Col=11 Row=G, DNA sequence.

ACCESSION AQ235963  
VERSION AQ235963.1 GI:3664570  
KEYWORDS GSS.  
SOURCE human.

## ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

## REFERENCE

## AUTHORS

1 (bases 1 to 516)  
Mahairas,G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,  
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and  
Hood,L.

## TITLE

Sequence-tagged connectors: A sequence approach to mapping and  
scanning the human genome

## JOURNAL

## MEDLINE

## COMMENT

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

Contact: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Sequence Tagged Connector

Plate: 2056 row: G column: 11

Class: BAC ends

High quality sequence stop: 516.

Location/Qualifiers  
1..516

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="Plate=2056 Col=11 Row=G"

/clone\_lib="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBelOBAC11; BAC Clones in  
E-Coli DH10B"

BASE COUNT 149 a 100 c 92 g 168 t 7 others  
ORIGIN

Query Match 71.2%; Score 17.8; DB 17; Length 516;

Best Local Similarity 82.6%; Pred. No. 2.2e+03;

Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CGCGTCGGTGCAGGCGTCACAA 23  
Db 27 CGCGTCGGTGCAGGTCACAA 5

RESULT 6

CNS04KEP/c

LOCUS 865 bp DNA linear GSS 21-MAY-2000

DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone  
116114 of library G from Tetraodon nigroviridis, genomic survey  
sequence.

AL294720

VERSION GSS: genome survey sequence.

KEYWORDS Tetraodon nigroviridis.

SOURCE Tetraodon nigroviridis.

ORGANISM Tetraodon nigroviridis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;

Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;

Tetraodontidae; Tetraodon.

1 (bases 1 to 865)

Roest-Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,

Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,

Saurin,W. and Weissenbach,J.

Human gene number estimate provided by genome wide analysis using  
Tetraodon nigroviridis DNA sequence

Unpublished

2 (bases 1 to 865)

Roest-Crolius,H., Jaillon,O., Dasilva,C., Fizames,C., Fisher,C.,

Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and

Weissenbach,J.

Characterization and repeat analysis of the compact genome of the  
freshwater pufferfish Tetraodon nigroviridis

Unpublished

3 (bases 1 to 865)

Genoscope.

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## REFERENCE

## AUTHORS

	Query Match	71.28;	Score 17.8;	DB 17;	Length 1031;	
	Best Local Similarity	90.5%;	Pred. No. 2.4e+03;	Mismatches 0;	Gaps 0;	
	Matches	19;	Conservative	0;	Mismatches 2;	Indels 0;
Oy	1	COCGTCGGTGCAGGACGTGC	21			
Dd	665	COCGTCGGAGCAGCGGAC	645			
	RESULT 8					
	LOCUS	BE190622				
	DEFINITION	S020d12.v1 Gm-cl037 Glycine max cDNA clone GENOME SYSTEMS CLONE ID:				
		Gm-cl037-2736 5', mRNA sequence.				
	ACCESSION	BE190622				
	VERSION	BE190622.1	GI:8669515			
	KEYWORDS	EST.				
	SOURCE	soybean.				
	ORGANISM	Glycine max				
		Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae; Glycine.				
	REFERENCE	1 (bases 1 to 199)				
	AUTHORS	Shoenmaker,R., Keim,P., Vodkin,L., Erpelting,J., Coryell,V., Khanna,A., Bolla,B., Marra,M., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T., Underwood,K., Steptoe,M., Theising,B., Allen,M., Bowers,Y., Person,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R., Waterston,R. and Willson,R.				
	TITLE	Public Soybean EST Project				
	JOURNAL	Unpublished (1999)				
	COMMENT	Contact: Shoenaker R/Public Soybean EST Project Public Soybean EST Project Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu Putative full length read vector to vector length is 200 This clone is available through: resGen, Invitrogen Corp. 2130 South Memorial Parkway Huntsville, AL 35801 For further information call: (800)-533-4363 or contact via email: cduresgen.com Insert Length: 817 Std Error: 0.00.				
	FEATURES					
		Location/Qualifiers				
		1..199				
		/organism="Glycine max"				
		/db_xref="taxon:3847"				
		/clone="GENOME SYSTEMS CLONE ID: Gm-cl037-2736"				
		/clone_lib="Gm-cl037"				
		/tissue_type="fully expanded leaves of greenhouse grown plants"				
		/dev_stage="2 week old"				
		/lab_host="DH10B"				
		/note="Vector: pSPORT1; Site_1: NotI; Site_2: SalI; This cDNA library was constructed from mRNA isolated from fully expanded leaves of greenhouse grown plants that were 2 weeks old. The library was prepared using the Life Technologies pSuperScript cDNA library construction kit. Complementary DNA was synthesized from mRNA using a poly(dT) sequence with a NotI restriction site. SalI linkers adapters were ligated to the blunt-ended cDNA fragments followed by NotI digestion. The cDNA fragments were directionally cloned into the NotI-SalI restriction site of the pSPORT1 vector. The ligated cDNA fragments were transformed into E.coli Electro-Max DH10B host cells. This library was constructed in the laboratory of Dr. Liia Vodkin by Anu Khanna at the University of Illinois at Urbana-Champaign. email: l-vodkin@uiuc.edu"				
		39 a	35 c	49 g	76 t	
	BASE COUNT					
	ORIGIN					
	Query Match	71.28;	Score 17.8;	DB 17;	Length 865;	
	Best Local Similarity	90.5%;	Pred. No. 2.4e+03;	Mismatches 0;	Gaps 0;	
	Matches	19;	Conservative	0;	Mismatches 2;	Indels 0;
Oy	1	COCGTCGGTGCAGGACGTGC	21			
Dd	235	COCGTCGGAGCAGCGGAC	215			
	RESULT 7					
	LOCUS	CNS04F4W/c				
	DEFINITION	Tetraodon nigroviridis genome survey sequence PUC-Ori end of clone 105J24 of library G from Tetraodon nigroviridis, genomic survey sequence.				
	ACCESSION	AL287897				
	VERSION	AL287897.1	GI:8026407			
	KEYWORDS	GSS; genome survey sequence.				
	SOURCE	Tetraodon nigroviridis.				
	ORGANISM	Tetraodon nigroviridis				
		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthomorpha; Acanthopterygii; Percormorpha; Tetraodontiformes; Tetraodontidae; Tetraodon.				
	REFERENCE	1 (bases 1 to 1031)				
	AUTHORS	Roest-Crollius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C., Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F., Saurin,W. and Weissenbach,J.				
	TITLE	Human gene number estimate provided by genome wide analysis using Tetraodon nigroviridis DNA sequence				
	JOURNAL	Unpublished				
	REFERENCE	2 (bases 1 to 1031)				
	AUTHORS	Roest-Crollius,H., Jaillon,O., Dasilva,C., Fizames,C., Fisher,C., Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and Weissenbach,J.				
	TITLE	Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetraodon nigroviridis				
	JOURNAL	Unpublished				
	REFERENCE	3 (bases 1 to 1031)				
	AUTHORS	Genoscope.				
	TITLE	Direct Submission				
	JOURNAL	Submitted (12-APR-2000)				
	COMMENT	This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetraodon nigroviridis genome. For more information, please take a look at http://www.genoscope.cns.fr/Tetraodon.				
	FEATURES					
		Location/Qualifiers				
		1..1031				
		/organism="Tetraodon nigroviridis"				
		/db_xref="taxon:99883"				
		/clone="105J24"				
		/clone_lib="G"				
		/note="Genoscope sequence ID : COBG105DE12SP1-end : PUC-Ori"				
	BASE COUNT	219 a	282 c			



Query Match 70.4%; Score 17.6; DB 10; Length 199;  
Best Local Similarity 83.3%; Pred. No. 2.2e+03;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CGCGTCGGTGCAGGACGTGACAAA 24  
||||| ||| | ||| ||||| |||  
Db 4 CGCGTGGGTCCTGGAGGTGACAAA 27

## RESULT 9

A0647334/c

## LOCUS

DEFINITION RPCI93-ECORI-2B12-TV RPCI93-ECORI Trypanosoma brucei genomic clone  
A0647334  
VERSION A0647334.1 GI:5140520  
KEYWORDS  
SOURCE Trypanosoma brucei.  
ORGANISM Trypanosoma brucei.

## REFERENCE

AUTHORS El-Sayed, N., Zhao, S., Zhao, H., Gill, S., Suh, E., Malek, J., Fujii, C., Gerrard, C., Leech, V., de Jong, P., Ullu, E., Melville, S., Donelson, J., Fraser, C. and Adams, M.  
TITLE Use of BAC end sequences from Trypanosoma brucei GUTat 10.1 RPCI-93  
JOURNAL Library for gene discovery and sequence-ready map construction  
COMMENT Other\_GSSs: RPCI93-ECORI-2B12.TP  
Contact: Najib M. El-Sayed  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: nelsayed@tigr.org  
Clones and high density filters may be purchased from BACPAC  
Resources (http://bacpac.med.buffalo.edu). BAC end sequences search  
page: http://www.tigr.org/tdb/mbd/tbdb/.

## FEATURES

source

1. .412  
/organism="Trypanosoma brucei"  
/strain="TREU927/4 GUTat 10.1"  
/db\_xref="taxon:5691"  
/clone="RPCI93-ECORI-2B12"  
/clone\_lib="RPCI93-ECORI"  
/note="Vector: pBACe3.6; Site\_1: Eco RI; Site\_2: Eco RI;  
Constructed for The Institute for Genomic Research by  
Bohui Zhao in Pieter de Jong's laboratory (Roswell Park  
Cancer Institute, Buffalo, NY). Briefly, Trypanosoma  
brucei TREU927/4 GUTat 10.1 agarose embedded DNA was  
partially digested with a combination of Eco RI and Eco RI  
methylase (RPCI93-ECORI segment) or Dpn II (RPCI93-DpnII  
segment). High molecular weight fragments were ligated in  
pBACe3.6 vector digested with Eco RI or Bam HI,  
respectively. The average insert size is 141 Kb. Total  
coverage (both segments): > 90 x the haploid  
non-minichromosomal genome." 95 g 97 t

## BASE COUNT

ORIGIN

Query Match 70.4%; Score 17.6; DB 17; Length 412;  
Best Local Similarity 83.3%; Pred. No. 2.5e+03;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 1 CGCGTCGGTGCAGGACGTGACAAA 24  
||||| ||| | ||| ||||| |||  
Db 231 CGCGCCAGTGGAGGAGGTGAAAA 208

## RESULT 10

A0641517/c

## LOCUS

## DEFINITION

RPCI93-ECORI-4M9-TV RPCI93-ECORI Trypanosoma brucei genomic clone

A0641517

## VERSION

A0641517.1 GI:5118227

## KEYWORDS

GSS.

## SOURCE

Trypanosoma brucei.

## ORGANISM

Trypanosoma brucei.

## REFERENCE

AUTHORS

1 (bases 1 to 414)

El-Sayed, N., Zhao, S., Zhao, H., Gill, S., Suh, E., Malek, J., Fujii, C., Gerrard, C., Leech, V., de Jong, P., Ullu, E., Melville, S., Donelson, J., Fraser, C. and Adams, M.

## TITLE

Library for gene discovery and sequence-ready map construction

## JOURNAL

COMMENT

Other\_GSSs: RPCI93-ECORI-4M9.TJ  
Contact: Najib M. El-Sayed  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: nelsayed@tigr.org  
Clones and high density filters may be purchased from BACPAC  
Resources (http://bacpac.med.buffalo.edu). BAC end sequences search  
page: http://www.tigr.org/tdb/mbd/tbdb/.

## FEATURES

source

1. .414

/organism="Trypanosoma brucei"  
/strain="TREU927/4 GUTat 10.1"  
/db\_xref="taxon:5691"  
/clone="RPCI93-ECORI-4M9"  
/clone\_lib="RPCI93-ECORI"  
/note="Vector: pBACe3.6; Site\_1: Eco RI; Site\_2: Eco RI;  
Constructed for The Institute for Genomic Research by  
Bohui Zhao in Pieter de Jong's laboratory (Roswell Park  
Cancer Institute, Buffalo, NY). Briefly, Trypanosoma  
brucei TREU927/4 GUTat 10.1 agarose embedded DNA was  
partially digested with a combination of Eco RI and Eco RI  
methylase (RPCI93-ECORI segment) or Dpn II (RPCI93-DpnII  
segment). High molecular weight fragments were ligated in  
pBACe3.6 vector digested with Eco RI or Bam HI,  
respectively. The average insert size is 141 Kb. Total  
coverage (both segments): > 90 x the haploid  
non-minichromosomal genome." 122 c 97 g 90 t

## BASE COUNT

ORIGIN

Query Match 70.4%; Score 17.6; DB 17; Length 414;  
Best Local Similarity 83.3%; Pred. No. 2.5e+03;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 1 CGCGTCGGTGCAGGACGTGACAAA 24  
||||| ||| | ||| ||||| |||  
Db 184 CGCGCCAGTGGAGGAGGTGAAAA 161

## RESULT 11

A2247056

## LOCUS

## DEFINITION

RPCI-23-41E10-TV RPCI-23 Mus musculus genomic clone RPCI-23-41E10,

A2247056

## ACCESSION

A2247056

## VERSION

A2247056.1 GI:8560271

## KEYWORDS

GSS.

## SOURCE

house mouse.

## ORGANISM

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al.  
 2001)  
 JOURNAL  
 COMMENT  
 Unpublished (2001)  
 Contact: Yoshihide Hayashizaki  
 Laboratory for Genome Exploration Research Group, RIKEN Genomic  
 Sciences Center(GSC), Yokohama Institute  
 The Institute of Physical and Chemical Research (RIKEN)  
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
 Tel: 81-45-503-9222  
 Fax: 81-45-503-9216  
 Email: genome-res@gsc.riken.go.jp/  
 URL:http://genome.gsc.riken.go.jp/  
 Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K., Itoh  
 M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.  
 Normalization and subtraction of cap-trapper-selected cDNAs to  
 prepare full-length cDNA libraries for rapid discovery of new  
 genes. Genome Res. 10 (10), 1617-1630 (2000)  
 wagi,K., Fujiwaka,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E.,  
 Watahiki,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsuura  
 S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and  
 Hayashizaki,Y.  
 RIKEN integrated sequence analysis (RISA) system--384-format  
 sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10  
 (11), 1757-1771 (2000)  
 Konno,H., Fukumishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara  
 Y., and Hayashizaki,Y.  
 Computer-based methods for the mouse full-length cDNA  
 encyclopedia: real-time sequence clustering for construction of a  
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)  
 Please visit our web site (http://genome.gsc.riken.go.jp) for  
 further details.  
 e mouse tissues.  
 Location/Qualifiers  
 1. 449  
 /organism="Mus musculus"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="E860105116"  
 /clone\_lib="RIKEN full-length enriched, 8 cells embryo"  
 /cell\_type="8 cells"  
 /dev\_stage="8 cells embryo"  
 /note="Vector: pSPOR1; Site\_1: SalI; Site\_2: NotI; This  
 clone is among a rearranged set of 15,247 clones from 11  
 embryo cDNA libraries (including preimplantation stage  
 embryos from unfertilized egg to blastocyst, embryonic  
 part of E7.5 embryos, extraembryonic part of E7.5 embryos  
 , and E12.5 female mesonephros/gonad) and one newborn  
 ovary cDNA library. Average insert size 1.5 kb. All  
 source libraries are cloned unidirectionally with Oligo(dT)  
 )-Not primers. References include: (1) Genome-wide  
 expression profiling of mid-gestation placenta and embryo  
 using a 15,000 mouse developmental cDNA microarray, 2000,  
 Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2)  
 Large-scale cDNA analysis reveals phased gene expression  
 patterns during preimplantation mouse development, 2000,  
 Development, 127: 1737-1749; (3) Genome-wide mapping of  
 unselected transcripts from extraembryonic tissue of  
 7.5-day mouse embryos reveals enrichment in the t-complex  
 and under-representation on the X chromosome, 1998, Hum  
 Mol Genet 7: 1967-1978."  
 144 a 95 c 80 g 130 t  
 -----  
 BASE COUNT  
 144 a 95 c 80 g 130 t  
 ORIGIN  
 Query Match 70.4%; Score 17.6; DB 10; Length 449;  
 Best Local Similarity 83.3%; Pred.No. 2 6e+03;  
 Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 2 GCGTCGGTGCAGGACGTGACAAAT 25  
 T TTT TTTTTTTTTTTTTTTTTTTTTT  
 Db 378 GGGTCTGTGCAGGACTTGACACAT 355  
 -----  
 RESULT 13

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 442)  
Zhao,S., Nierman,W., Feldblyum,T., Malek,J., Shatsman,S., Akinret,  
B., Levins,M., McGann,S., Tsegaye,G., Geer,K., Krol,M., de Jong,P.  
and Fraser,C.M.  
Mouse BAC End Sequences from Library RPCI-23  
Unpublished (1999)  
Other\_GSSs: RPCI-23-41E10.TJ  
Contact: Shaying Zhao  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: szhao@tigr.org  
Clones are derived from the mouse BAC library RPCI-23. For BAC  
library availability, please contact Pieter de Jong  
(pieterdejong.med.buffalo.edu). Clones may be purchased from  
BACPAC Resources (<http://bacpac.med.buffalo.edu/orderingframe.htm>)  
or from Resea ch Genetics ([inforesgen.com](http://inforesgen.com)). BAC end page:  
[http://www.tigr.org/tdb/bac\\_ends/mouse/bac\\_end\\_intro.html](http://www.tigr.org/tdb/bac_ends/mouse/bac_end_intro.html)  
Plate: 41 row: E Column: 10  
Seq primer: F7  
Class: BAC ends.  
Location/Qualifiers  
1. .442  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="RPCI-23-41E10"  
/clone\_lib="RPCI-23"  
/sex="Female"  
/lab\_host="DH10B"  
/note="Organ: Kidney/Brain; Vector: pBACe3.6; Site:1:  
EcoRI; Site:2: EcoRI; Female C57BL/6J mouse kidney and/or  
brain genomic DNA was isolated and partially digested  
with a combination of EcoRI and EcoRI Methylase. Size  
selected DNA was cloned into the pBACe3.6 vector at the  
EcoRI sites. The ligation products were transformed into  
DH10B electrocompetent cells (BRL Life Technologies)."  
BASE COUNT  
128 a 82 c 89 g 143 t  
ORIGIN  
Query Match 70.4%; Score 17.6; DB 17; Length 442;  
Best Local Similarity 83.3%; Pred. No. 2 5e-03;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 2 GCGTCGGTGCAGGCGTGCACAAAT 25  
||||| ||||||| ||||| ||  
Db 83 GGGTGTGTGCAGGACTTGACACAT 106  
RESULT 12  
BB839535/c  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
house mouse.  
Mus musculus  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE  
AUTHORS  
Akimura,T., Arakawa,T., Carninci,P., Furuno,M., Hanagaki,T.,  
Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,T., Imotani,K., Ishii,  
Y., Ito,M., Kawai,J., Kojima,Y., Konno,H., Konda,M., Matsuyama,T.,  
Nakamura,M., Nishi,K., Nomura,K., Numasaki,R., Okazaki,Y., Okido,T.,  
Saito,R., Sakai,C., Sakai,K., Sakazume,N., Sasaki,D., Sato,K.,  
Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa,  
A., Takahashi,F., Takaku-Akahira,S., Tanaka,T., Tomaru,A., Toya,T.,  
Uwatashi,A., Yasunishi,A., Muramatsu,M. and Hayashizaki,Y.

AQ844401  
LOCUS an36h09 JM101 filtered library Zea mays genomic, DNA sequence.  
ACCESSION AQ844401  
VERSION AQ844401.1 GI:6202889  
KEYWORDS GSS.  
SOURCE Zea mays.  
ORGANISM Zea mays.  
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC clade; Panicoideae; Andropogoneae; Zea.  
AUTHORS 1 (bases 1 to 464)  
Rabinowicz,P.D., Schutz,K., Dedhia,N., Yordan,C., Parnell,L.D., Stein,L., McCombie,W.R. and Martienssen,R.A.  
TITLE Differential methylation of genes and retrotransposons allows shotgun sequencing of the maize genome  
JOURNAL Nat. Genet. 23, 305-308 (1999)  
COMMENT Contact: Martienssen RA  
Cold Spring Harbor Laboratory  
1 Bungtown Rd., Cold Spring Harbor, NY 11724, USA  
Tel: 516 367 8322  
Fax: 516 367 8369  
Email: martiensscshl.org  
Seq primer: forward  
Class: Shotgun.  
FEATURES  
source  
1. 464  
/organism="Zea mays"  
/cultivar="B73"  
/db\_xref="taxon:4577"  
/clone\_lib="JM101 filtered library"  
/note="Organ: immature ears; Vector: M13; Site\_1: Xba I; DNA prepared from purified nuclei was digested with the methylation insensitive enzyme Spe I, size fractionated to enrich for the 0.5 to 4 kbp fraction, ligated into Xba I digested M13 vector and electroporated into E.coli JM101."  
BASE COUNT 77 a 146 c 156 g 85 t  
ORIGIN  
Query Match 70.4%; Score 17.6; DB 17; Length 464;  
Best Local Similarity 83.3%; Pred. No. 2.6e+03;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 1 CGCGTCGTCGACGACGTGCATA 24  
||||| ||||||||| |||  
Db 149 CGCGTCGTCGACGACGTGCATA 172  
RESULT 14  
AA957358 515 bp mRNA linear EST 04-JUL-1999  
LOCUS UI-R-EI-fy-b-07-0-UI-s1 UI-R-EI Rattus norvegicus cDNA clone  
DEFINITION UI-R-EI-fy-b-07-0-UI 3', mRNA sequence.  
ACCESSION AA957358  
VERSION AA957358.1 GI:4277248  
KEYWORDS EST.  
SOURCE Norway rat.  
ORGANISM Rattus norvegicus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
REFERENCE 1 (bases 1 to 515)  
AUTHORS Ronaldo,M.F., Lennon,G. and Soares,M.B.  
TITLE Normalization and subtraction: two approaches to facilitate gene discovery  
JOURNAL Genome Res. 6 (9), 791-806 (1996)  
MEDLINE 97044477  
COMMENT On May 7, 1998 this sequence version replaced gi:3121053.  
Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA

Tel: 319 335 8250  
Fax: 319 335 9565  
Email: msoares@blue.weeg.uiowa.edu  
Oligo-dT track not found, Not 1 site shown in beginning of sequence.  
is likely internal to the message. cDNA library preparation: M.  
Fatima Bonaldo, Ph.D. Clone distribution: clones will be available  
through Research Genetics This clone is also available through the  
I.M.A.G.E. Consortium at LLNL (info@image.llnl.gov). IMAGE  
ID=1780190 The following repetitive elements were found in this  
cDNA sequence: 57-96, >GC-richLow\_complexity  
Seq primer: M13 Forward  
POLYA-No. Location/Qualifiers  
1. 515  
/organism="Rattus norvegicus"  
/strain="Sprague-Dawley"  
/db\_xref="taxon:10116"  
/clone="UI-R-EI-fy-b-07-0-UI"  
/clone\_lib="UI-R-EI"  
/dev\_stage="adult"  
/lab\_host="DH10B (Life Technologies)"  
/note="Vector: pT73D-pac (Pharmacia) with a modified  
polylinker: Site\_1: Not I; Site\_2: Eco RI; The UI-R-EI  
library is a subtracted library derived from the UI-R-EI  
library. The UI-R-EI library consisted of a mixture of  
individually tagged normalized libraries constructed from  
8, 12 and 18-day embryo. The tag is a string of 3-5  
nucleotides present between the Not I site and the  
oligo-dT track which allows identification of the library  
of origin of a clone within the mixture. The subtracted  
library (UI-R-EI) was constructed as follows: PCR  
amplified cDNA inserts from a pool of UI-R-EI clones from  
which 3' ESTs had been derived was used as a driver in a  
hybridization with the UI-R-EI library in the form of  
single-stranded circles. The remaining single-stranded  
circles (subtracted library) was purified by  
hydroxyapatite column chromatography, converted to  
double-stranded circles and electroporated into DH10B  
bacteria (Life Technologies) to generate the UI-R-EI  
library. This procedure has been previously described  
(Bonaldo, Lennon and Soares, Genome Research 6: 791-806,  
1996)."  
BASE COUNT 159 a 116 c 149 g 91 t  
ORIGIN  
Query Match 70.4%; Score 17.6; DB 9; Length 515;  
Best Local Similarity 83.3%; Pred. No. 2.6e+03;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 1 CGCGTCGTCGACGACGTGCACAA 24  
||||| ||||||||| |||  
Db 467 CACGTCGTCGACGACGTGGCAAA 490  
RESULT 15  
AW664262/c 564 bp mRNA linear EST 06-APR-2000  
LOCUS h108e10.x1 NCI\_CGAP\_GUI Homo sapiens cDNA clone IMAGE:2971722 3'  
DEFINITION Similar to FR:O35540 O35540 HEPATOMA-DERIVED GROWTH FACTOR, RELATED  
PROTEIN 2. ; mRNA sequence.  
ACCESSION AW664262  
VERSION AW664262.1 GI:7456803  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 564)  
AUTHORS NCI\_CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index  
JOURNAL Unpublished (1997)  
COMMENT Contact: Robert Strausberg, Ph.D.

Mon Jan 6 15:20:17 2003

Email: cgapbs-re@mail.nih.gov  
Tissue Procurement: Chris Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D. cDNA Library Preparation: Life Technologies, Inc. cDNA Library Arrayed by: Christa Prange, The I.M.A.G.E. Consortium DNA Sequencing by: Washington University Genome Sequencing Center  
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov/image/html/iresources.shtml

Possible reversed clone: polyt not found

Seq primer: -40UP from Gibco

High quality sequence stop: 398.

# FEATURES

source

Location/Qualifiers

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/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="IMAGE:2971722"

/clone\_lib="NCI-CGAP\_GUI"

/tissue\_type="2 pooled high-grade transitional cell

tumors"

/lab\_host="DH10B"

/note="Organ: genitourinary tract; Vector: pCMV-SPORT6;

Site\_1: SalI; Site\_2: NotI; Cloned unidirectionally.

Primer: Oligo dT. Library constructed by Life

Technologies."

BASE COUNT 85 a 202 c 134 g 142 t 1 others

ORIGIN

Query Match

Best Local Similarity 70.4%; Score 17.6; DB 10; Length 564;

Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

0;

Qy 1 CGCGTCGGTGCAGGACGTGACAAA 24

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Db 562 CGAGCGGTGCAGAAAGTGACAAA 539

Search completed: January 4, 2003, 01:04:07

Job time : 338.413 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:53:16 ; Search time 8.43849 Seconds  
(without alignments)  
908.566 Million cell updates/sec

Title: US-09-787-562-1

Perfect score: 25

Sequence: 1 cgcgcggtgcaggacgtgacaaat 25

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued\_Patents\_NA:\*

1: /cgn2\_6/ptodata/2/ina/5A.COMB.seq.\*

2: /cgn2\_6/ptodata/2/ina/5B.COMB.seq.\*

3: /cgn2\_6/ptodata/2/ina/6A.COMB.seq.\*

4: /cgn2\_6/ptodata/2/ina/6B.COMB.seq.\*

5: /cgn2\_6/ptodata/2/ina/PCTUS.COMB.seq.\*

6: /cgn2\_6/ptodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

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2	18.6	74.4	3670	2	US-08-600-452A-3
3	18	72.0	24	2	US-08-693-174-2
4	18	72.0	24	4	US-09-253-738-2
5	18	72.0	1110	2	US-08-693-174-4
6	18	72.0	1110	4	US-09-253-738-4
7	18	72.0	5382	4	US-09-479-122-21
8	18	72.0	7617	3	US-08-646-538-34
9	18	72.0	7617	3	US-09-503-222-34
10	18	72.0	8387	2	US-08-532-814-1
11	18	72.0	8388	4	US-09-225-509-1
12	18	72.0	9737	4	US-09-479-122-22
13	18	72.0	9737	4	US-09-479-122-23
14	18	72.0	9737	4	US-09-479-122-28
15	18	72.0	9871	4	US-09-479-122-28
16	18	72.0	10060	4	US-09-479-122-25
17	17.6	70.4	4256	1	US-08-505-509-31
18	17.6	70.4	4256	2	US-08-491-690A-31
19	16.2	64.8	6671	1	US-08-280-443-1
20	16.2	64.8	6671	1	US-08-457-459-1
21	16.2	64.8	6671	1	US-08-555-678-1
22	16.2	64.8	6671	5	PCT-US95-02275-1
23	16	64.0	1464	4	US-09-351-224E-4
24	16	64.0	2353	5	PCT-US92-06840-1
25	16	64.0	3735	4	US-08-975-762-43
26	16	64.0	3735	4	US-09-295-028-43
27	16	64.0	3735	4	US-09-106-582-43

28 16 64.0 12847 1 US-08-550-715-1 Sequence 1, Appli

29 16 64.0 4411529 4 US-09-103-840A-1 Sequence 1, Appli

30 15.8 63.2 372 4 US-09-124-671-24 Sequence 24, Appli

31 15.8 63.2 1815 3 US-09-041-545-1 Sequence 1, Appli

32 15.8 63.2 1815 3 US-09-327-925-1 Sequence 1, Appli

33 15.8 63.2 18627 4 US-08-961-527-113 Sequence 113, App

34 15.8 63.2 43950 4 US-09-735-934A-3 Sequence 3, Appli

35 15.8 63.2 4403765 4 US-09-103-840A-2 Sequence 2, Appli

36 15.6 62.4 200 1 US-08-308-892A-13 Sequence 13, Appli

37 15.6 62.4 200 1 US-08-308-892A-14 Sequence 14, Appli

38 15.6 62.4 200 1 US-08-308-892A-15 Sequence 15, Appli

39 15.6 62.4 274 4 US-08-990-823-72 Sequence 72, Appli

40 15.6 62.4 507 1 US-08-246-420A-1 Sequence 1, Appli

41 15.6 62.4 507 2 US-08-766-620A-1 Sequence 1, Appli

42 15.6 62.4 507 5 PCT-US95-06094-1 Sequence 1, Appli

43 15.6 62.4 525 1 US-08-009-973-2 Sequence 2, Appli

44 15.6 62.4 540 4 US-08-149-101A-3 Sequence 3, Appli

45 15.6 62.4 540 5 PCT-US94-12873-3 Sequence 3, Appli

## ALIGNMENTS

RESULT 1

US-08-386-727-3

; Sequence 3, Application US/08386727

; Patent No. 5792647

; GENERAL INFORMATION:

; APPLICANT: ROSEMAN, SAUL

; APPLICANT: BASSLER, BONNIE

; APPLICANT: KEYHANT, NEMAT O.

; APPLICANT: CHITLARU, EDITH

; APPLICANT: ROWE, CHRIS

; APPLICANT: YU, CHARLES

; TITLE OF INVENTION: BACTERIAL CATABOLISM OF CHITIN

; NUMBER OF SEQUENCES: 8

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: CUSHMAN, DARBY & CUSHMAN

; STREET: 1100 NEW YORK AVENUE, N.W.

; CITY: WASHINGTON

; STATE: D.C.

; COUNTRY: USA

; ZIP: 20005

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/386,727

; FILING DATE:

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: HOBBS, ANN S.

; REGISTRATION NUMBER: 36,830

; REFERENCE/DOCKET NUMBER: 4130/206916

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 202-861-3000

; TELEX: 6714627 CUSH

; INFORMATION FOR SEQ ID NO: 3:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 3670 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: DNA (genomic)

US-08-386-727-3

Query Match 74.4%; Score 18.6; DB 1; Length 3670;

Best Local Similarity 84.0%; Pred. No. 7.9;

Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;



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; OTHER INFORMATION: a, c, t, g, other or unknown
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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      8 GTGCAGGACGTGACAAAT 25
Db      899 GTGCAGGACGTGACAAAT 916

RESULT 8
US-08-646-538-34/C
; Sequence 34, Application US/08646538
; Patent No. 6027881
; GENERAL INFORMATION:
; APPLICANT: Pavlakis, George N.
; APPLICANT: Gaitanaris, George A.
; APPLICANT: Stauber, Roland H.
; APPLICANT: Vournakis, John N.
; TITLE OF INVENTION: Mutant Aequorea victoria Fluorescent
; NUMBER OF SEQUENCES: 37
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, 8th Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC Compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/646,538
; FILING DATE: No. 6027881 yet assigned
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weber, Kenneth A.
; REGISTRATION NUMBER: 31,677
; REFERENCE/DOCKET NUMBER: 015280-249000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 34:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7617 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; FEATURE:
; NAME/KEY: -
; LOCATION: 1..7617
; OTHER INFORMATION: /note= "pGen-PGk9fo25RO"
US-08-646-538-34

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Qy      8 GTGCAGGACGTGACAAAT 25
Db      4186 GTGCAGGACGTGACAAAT 4169

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; Sequence 34, Application US/09503222
; Patent No. 6265548

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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      8 GTGCAGGACGTGACAAAT 25
Db      637 GTGCAGGACGTGACAAAT 654

RESULT 6
US-09-253-738-4
; Sequence 4, Application US/09253738
; Patent No. 6265390
; GENERAL INFORMATION:
; APPLICANT: Ratcliffe, Peter John
; APPLICANT: Firth, John David
; APPLICANT: Harris, Adrian Llewellyn
; APPLICANT: Pugh, Christopher William
; APPLICANT: Stratford, Ian James
; TITLE OF INVENTION: Targeting Gene Therapy
; FILE REFERENCE: 08/693174
; CURRENT APPLICATION NUMBER: US/09/253,738
; CURRENT FILING DATE: 1999-02-22
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 1110
; TYPE: DNA
; ORGANISM: Murinae gen. sp.
US-09-253-738-4

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Best Local Similarity 100.0%; Pred. No. 13;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      8 GTGCAGGACGTGACAAAT 25
Db      637 GTGCAGGACGTGACAAAT 654

RESULT 7
US-09-479-122-21
; Sequence 21, Application US/09479122
; Patent No. 6410266
; GENERAL INFORMATION:
; APPLICANT: HARRINGTON, JOHN J.
; APPLICANT: SHERF, BRUCE
; APPLICANT: RUNDLETT, STEPHEN
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR NON-TARGETED ACTIVATION OF
; FILE OF INVENTION: ENDOGENOUS GENES
; FILE REFERENCE: 0221-0003C
; CURRENT APPLICATION NUMBER: US/09/479,122
; CURRENT FILING DATE: 2000-01-07
; PRIOR APPLICATION NUMBER: 09/276,820
; PRIOR FILING DATE: 1999-03-26
; PRIOR APPLICATION NUMBER: 09/159,643
; PRIOR FILING DATE: 1998-09-24
; PRIOR APPLICATION NUMBER: 08/941,223
; PRIOR FILING DATE: 1997-09-26
; PRIOR APPLICATION NUMBER: 09/263,814
; PRIOR FILING DATE: 1999-03-08
; PRIOR APPLICATION NUMBER: 09/253,022
; PRIOR FILING DATE: 1999-02-19
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 21
; LENGTH: 5382
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (890)
; OTHER INFORMATION: a, c, t, g, other or unknown
; NAME/KEY: modified_base
; LOCATION: (1042)

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; GENERAL INFORMATION:
; APPLICANT: Pavlakis, George N.
; APPLICANT: Gaitanaris, George A.
; APPLICANT: Stauber, Roland H.
; APPLICANT: Vournakis, John N.
; TITLE OF INVENTION: Mutant Aequorea victoria Fluorescent
; TITLE OF INVENTION: Proteins Having Increased Cellular Fluorescence
; NUMBER OF SEQUENCES: 37
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, 8th Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/503,222
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/646,538
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Weber, Kenneth A.
; REGISTRATION NUMBER: 31,677
; REFERENCE/DOCKET NUMBER: 015280-249000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 34:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7617 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; FEATURE:
; NAME/KEY: -
; LOCATION: 1..7617
; OTHER INFORMATION: /note= "pGen-pkgfo25RO"
; US-09-503-222-34

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Best Local Similarity 100.0%; Pred. No. 16;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGAGCGTGACAAAT 25
Db 4186 GTGCAGGAGCGTGACAAAT 4169
|||||
; GENERAL INFORMATION:
; APPLICANT: MOULLIER, PHILIPPE
; APPLICANT: DANOS, OLIVIER
; APPLICANT: HEARD, JEAN-MICHEL
; APPLICANT: FERRY, NICOLAS
; TITLE OF INVENTION: BIOCOMPATIBLE IMPLANT FOR THE EXPRESSION
; TITLE OF INVENTION: AND IN VIVO SECRETION OF A THERAPEUTIC SUBSTANCE
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; STREET: P.C.
; STREET: 1755 S. JEFFERSON DAVIS HIGHWAY, SUITE 400
; CITY: ARLINGTON
; STATE: VIRGINIA
; COUNTRY: USA
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/532,814
; FILING DATE: 19-JAN-1996
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/FR94/00456
; FILING DATE: 21-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 93/04700
; FILING DATE: 21-APR-1993
; APPLICATION NUMBER: FR 93/09185
; FILING DATE: 26-JUL-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: OBLON, NORMAN F.
; REGISTRATION NUMBER: 24,618
; REFERENCE/DOCKET NUMBER: 660-105-0 PCT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 703-413-3000
; TELEFAX: 703-413-2220
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 8387 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-532-814-1

Query Match 72.0%; Score 18; DB 2; Length 8387;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGAGCGTGACAAAT 25
Db 2238 GTGCAGGAGCGTGACAAAT 2255
|||||
; GENERAL INFORMATION:
; APPLICANT: MOULLIER, Philippe
; APPLICANT: DANOS, Olivier
; APPLICANT: HEARD, Jean-Michel
; APPLICANT: FERRY, Nicholas
; TITLE OF INVENTION: BIOCOMPATIBLE IMPLANT FOR THE EXPRESSION AND IN VIVO
; TITLE OF INVENTION: SECRETION OF A THERAPEUTIC SUBSTANCE
; FILE REFERENCE: 0660-0145-0DIV
; CURRENT APPLICATION NUMBER: US/09/225,509
; CURRENT FILING DATE: 1999-01-06
; EARLIER APPLICATION NUMBER: PCT/FR94/00456
; EARLIER FILING DATE: 1994-04-21
; EARLIER APPLICATION NUMBER: 09/523,814
; EARLIER FILING DATE: 1996-01-19
; EARLIER APPLICATION NUMBER: FR 93/04700
; EARLIER FILING DATE: 1993-04-21
; EARLIER APPLICATION NUMBER: FR 93/09185
; EARLIER FILING DATE: 1993-07-26
; NUMBER OF SEQ ID NOS: 1
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 8388
; TYPE: DNA
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; US-09-225-509-1
; Sequence 1, Application US/09225509
; Patent No. 6326195
; GENERAL INFORMATION:
; APPLICANT: MOULLIER, Philippe
; APPLICANT: DANOS, Olivier
; APPLICANT: HEARD, Jean-Michel
; APPLICANT: FERRY, Nicholas
; TITLE OF INVENTION: BIOCOMPATIBLE IMPLANT FOR THE EXPRESSION AND IN VIVO
; TITLE OF INVENTION: SECRETION OF A THERAPEUTIC SUBSTANCE
; FILE REFERENCE: 0660-0145-0DIV
; CURRENT APPLICATION NUMBER: US/09/225,509
; CURRENT FILING DATE: 1999-01-06
; EARLIER APPLICATION NUMBER: PCT/FR94/00456
; EARLIER FILING DATE: 1994-04-21
; EARLIER APPLICATION NUMBER: 09/523,814
; EARLIER FILING DATE: 1996-01-19
; EARLIER APPLICATION NUMBER: FR 93/04700
; EARLIER FILING DATE: 1993-04-21
; EARLIER APPLICATION NUMBER: FR 93/09185
; EARLIER FILING DATE: 1993-07-26
; NUMBER OF SEQ ID NOS: 1
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 8388
; TYPE: DNA
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; US-09-225-509-1
; Sequence 1, Application US/09225509
; Patent No. 6326195
; GENERAL INFORMATION:
; APPLICANT: MOULLIER, Philippe
; APPLICANT: DANOS, Olivier
; APPLICANT: HEARD, Jean-Michel
; APPLICANT: FERRY, Nicholas
; TITLE OF INVENTION: BIOCOMPATIBLE IMPLANT FOR THE EXPRESSION AND IN VIVO
; TITLE OF INVENTION: SECRETION OF A THERAPEUTIC SUBSTANCE
; FILE REFERENCE: 0660-0145-0DIV
; CURRENT APPLICATION NUMBER: US/09/225,509
; CURRENT FILING DATE: 1999-01-06
; EARLIER APPLICATION NUMBER: PCT/FR94/00456
; EARLIER FILING DATE: 1994-04-21
; EARLIER APPLICATION NUMBER: 09/523,814
; EARLIER FILING DATE: 1996-01-19
; EARLIER APPLICATION NUMBER: FR 93/04700
; EARLIER FILING DATE: 1993-04-21
; EARLIER APPLICATION NUMBER: FR 93/09185
; EARLIER FILING DATE: 1993-07-26
; NUMBER OF SEQ ID NOS: 1
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 8388
; TYPE: DNA
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; ORGANISM: mus musculus, Mo-MuLV, and other
US-09-225-509-1
Query Match          72.0%; Score 18; DB 4; Length 8388;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 GTGCAGGACGTGACAAAT 25
    |||||
Db 2238 GTGCAGGACGTGACAAAT 2255

RESULT 12
US-09-479-122-22
; Sequence 22, Application US/09479122
; Patent No. 6410266
; GENERAL INFORMATION:
; APPLICANT: HARRINGTON, JOHN J.
; APPLICANT: SHERF, BRUCE
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR NON-TARGETED ACTIVATION OF
; FILE REFERENCE: 0221-0003C
; CURRENT APPLICATION NUMBER: US/09/479,122
; CURRENT FILING DATE: 2000-01-07
; PRIOR APPLICATION NUMBER: 09/276,820
; PRIOR FILING DATE: 1999-03-26
; PRIOR APPLICATION NUMBER: 09/159,643
; PRIOR FILING DATE: 1998-09-24
; PRIOR APPLICATION NUMBER: 08/941,223
; PRIOR FILING DATE: 1997-09-26
; PRIOR APPLICATION NUMBER: 09/263,814
; PRIOR FILING DATE: 1999-03-08
; PRIOR APPLICATION NUMBER: 09/253,022
; PRIOR FILING DATE: 1999-02-19
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 22
; LENGTH: 9737
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (8347)
; OTHER INFORMATION: a, c, t, g, other or unknown
; NAME/KEY: modified_base
; LOCATION: (8499)
; OTHER INFORMATION: a, c, t, g, other or unknown
US-09-479-122-23
Query Match          72.0%; Score 18; DB 4; Length 9737;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 GTGCAGGACGTGACAAAT 25
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Db 8356 GTGCAGGACGTGACAAAT 8373

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; Patent No. 6410266
; GENERAL INFORMATION:
; APPLICANT: HARRINGTON, JOHN J.
; APPLICANT: SHERF, BRUCE
; APPLICANT: RUNDLETT, STEPHEN
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR NON-TARGETED ACTIVATION OF
; FILE REFERENCE: 0221-0003C
; CURRENT APPLICATION NUMBER: US/09/479,122
; CURRENT FILING DATE: 2000-01-07
; PRIOR APPLICATION NUMBER: 09/276,820
; PRIOR FILING DATE: 1999-03-26
; PRIOR APPLICATION NUMBER: 09/159,643
; PRIOR FILING DATE: 1998-09-24
; PRIOR APPLICATION NUMBER: 08/941,223
; PRIOR FILING DATE: 1997-09-26
; PRIOR APPLICATION NUMBER: 09/263,814
; PRIOR FILING DATE: 1999-03-08
; PRIOR APPLICATION NUMBER: 09/253,022
; PRIOR FILING DATE: 1999-02-19
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 28
; LENGTH: 9737
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (8347)
; OTHER INFORMATION: a, c, t, g, other or unknown
; NAME/KEY: modified_base
; LOCATION: (8499)
; OTHER INFORMATION: a, c, t, g, other or unknown
US-09-479-122-28
Query Match          72.0%; Score 18; DB 4; Length 9737;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 8356 GTGCAGGACGTGACAAAT 8373

RESULT 13
US-09-479-122-23
; Sequence 23, Application US/09479122
; Patent No. 6410266
; GENERAL INFORMATION:
; APPLICANT: HARRINGTON, JOHN J.
; APPLICANT: SHERF, BRUCE
; APPLICANT: RUNDLETT, STEPHEN
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR NON-TARGETED ACTIVATION OF
; FILE REFERENCE: 0221-0003C
; CURRENT APPLICATION NUMBER: US/09/479,122
; CURRENT FILING DATE: 2000-01-07
; PRIOR APPLICATION NUMBER: 09/276,820
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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
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 Db 8356 GTGCAGGACGTGACAAAT 8373

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 US-09-479-122-24  
 ; Sequence 24, Application US/09479122  
 ; Patent No. 6410266  
 ; GENERAL INFORMATION:  
 ; APPLICANT: HARRINGTON, JOHN J.  
 ; APPLICANT: SHERF, BRUCE  
 ; APPLICANT: RUNDLETT, STEPHEN  
 ; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR NON-TARGETED ACTIVATION OF  
 ; TITLE OF INVENTION: ENDOGENOUS GENES  
 ; FILE REFERENCE: 0221-0003C  
 ; CURRENT APPLICATION NUMBER: US/09/479,122  
 ; CURRENT FILING DATE: 2000-01-07  
 ; PRIOR APPLICATION NUMBER: 09/276,820  
 ; PRIOR FILING DATE: 1999-03-26  
 ; PRIOR APPLICATION NUMBER: 09/159,643  
 ; PRIOR FILING DATE: 1998-09-24  
 ; PRIOR APPLICATION NUMBER: 08/941,223  
 ; PRIOR FILING DATE: 1997-09-26  
 ; PRIOR APPLICATION NUMBER: 09/263,814  
 ; PRIOR FILING DATE: 1999-03-08  
 ; PRIOR APPLICATION NUMBER: 09/253,022  
 ; PRIOR FILING DATE: 1999-02-19  
 ; NUMBER OF SEQ ID NOS: 33  
 ; SOFTWARE: PatentIn Ver. 2.1  
 ; SEQ ID NO 24  
 ; LENGTH: 9871  
 ; TYPE: DNA  
 ; ORGANISM: Homo sapiens  
 ; FEATURE:  
 ; NAME/KEY: modified\_base  
 ; LOCATION: (8481)  
 ; OTHER INFORMATION: a, c, t, g, other or unknown  
 ; NAME/KEY: modified\_base  
 ; LOCATION: (8633)  
 ; OTHER INFORMATION: a, c, t, g, other or unknown  
 US-09-479-122-24

Query Match 72.0%; Score 18; DB 4; Length 9871;  
 Best Local Similarity 100.0%; Pred. No. 17;  
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
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 Db 8490 GTGCAGGACGTGACAAAT 8507

Search completed: January 3, 2003, 23:53:38  
 Job time : 14.4385 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:41 ; Search time 8.43849 Seconds  
(without alignments)  
1281.345 Million cell updates/sec

Title: US-09-787-562-1

Perfect score: 25

Sequence: 1 cgcctcggtgcaggacgtgacaaat 25

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 381593 seqs, 216252194 residues

Total number of hits satisfying chosen parameters: 763186

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Published Applications.NA.\*  
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3: /cgn2\_6/ptodata/1/pubpna/US06\_NEW\_PUB.seq.\*  
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13: /cgn2\_6/ptodata/1/pubpna/US60\_NEW\_PUB.seq.\*  
14: /cgn2\_6/ptodata/1/pubpna/US60\_PUBCOMB.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	72.0	943	10	US-09-833-381-1259
2	18	72.0	4768	9	US-10-087-523-1
3	18	72.0	4768	10	US-09-816-790-1
4	18	72.0	4768	10	US-09-861-077-1
5	18	72.0	4768	10	US-09-815-825-1
6	18	72.0	4768	10	US-09-815-935-1
7	18	72.0	4768	10	US-09-815-944-1
8	18	72.0	6355	9	US-10-087-523-2
9	18	72.0	6355	10	US-09-816-790-2
10	18	72.0	6355	10	US-09-861-077-2
11	18	72.0	6355	10	US-09-815-825-2
12	18	72.0	6355	10	US-09-815-935-2
13	18	72.0	6355	10	US-09-815-944-2
14	18	72.0	8388	10	US-09-987-601-1
15	17	68.0	279	10	US-09-919-580-500
16	16.2	64.8	279	10	US-09-878-574-13168
17	16.2	64.8	538	10	US-09-998-598-458
18	16.2	64.8	729	9	US-09-738-626-1680
19	16.2	64.8	1797	9	US-09-738-626-1682

c	20	16	64.0	235	10	US-09-736-960-66	Sequence 66, Appl
c	21	16	64.0	534	9	US-09-738-626-2011	Sequence 2011, Ap
c	22	16	64.0	987	10	US-09-974-300-2046	Sequence 2046, Ap
c	23	16	64.0	1314	9	US-09-738-626-1619	Sequence 1619, Ap
c	24	16	64.0	1852	10	US-09-925-302-58	Sequence 58, Appl
c	25	16	64.0	3735	10	US-09-159-469-43	Sequence 43, Appl
c	26	16	64.0	3735	10	US-09-798-042-43	Sequence 43, Appl
c	27	16	64.0	3735	10	US-09-798-042-88	Sequence 88, Appl
c	28	16	64.0	3735	10	US-09-798-042-96	Sequence 96, Appl
c	29	16	64.0	4026	10	US-09-736-960-3	Sequence 3, Appl
c	30	16	64.0	7215	10	US-09-736-960-1	Sequence 1, Appl
c	31	16	64.0	43058	10	US-09-954-456-292	Sequence 292, App
c	32	16	64.0	43058	10	US-09-954-456-529	Sequence 529, App
c	33	16	64.0	43058	10	US-09-880-107-3950	Sequence 3950, Ap
c	34	16	64.0	66686	10	US-09-736-960-86	Sequence 86, Appl
c	35	16	64.0	3309400	9	US-09-738-626-1	Sequence 1, Appl
c	36	15.8	63.2	274	10	US-09-878-574-10968	Sequence 10968, A
c	37	15.8	63.2	461	10	US-09-974-300-8244	Sequence 8244, Ap
c	38	15.8	63.2	755	10	US-09-919-603-4	Sequence 4, Appl
c	39	15.8	63.2	925	10	US-09-919-603-6	Sequence 6, Appl
c	40	15.8	63.2	2439	9	US-09-954-531-140	Sequence 140, App
c	41	15.8	63.2	2439	9	US-09-954-531-359	Sequence 359, App
c	42	15.8	63.2	2748	10	US-09-822-849A-234	Sequence 234, App
c	43	15.8	63.2	43950	12	US-10-060-332-3	Sequence 3, Appl
c	44	15.6	62.4	274	9	US-09-996-634-72	Sequence 72, Appl
c	45	15.6	62.4	425	10	US-09-983-965-2423	Sequence 2423, Ap

#### ALIGNMENTS

##### RESULT 1

US-09-833-381-1259/c

; Sequence 1259, Application US/09833381

; Patent No. US20020132090A1

; GENERAL INFORMATION:

; APPLICANT: Robison, Keith E.

; TITLE OF INVENTION: No. US20020132090A1el Nucleic Acid and Protein Homologs

; FILE REFERENCE: 5800-119

; CURRENT APPLICATION NUMBER: US/09/833.381

; PRIOR FILING DATE: 2001-04-11

; PRIOR APPLICATION NUMBER: 09/516,448

; FILING DATE: 2000-02-29

; NUMBER OF SEQ ID NOS: 2050

; SOFTWARE: FastSeq for Windows Version 3.0

; SEQ ID NO 1259

; LENGTH: 943

; TYPE: DNA

; ORGANISM: Homo sapiens

; FEATURE:

; NAME/KEY: misc.feature

; LOCATION: (1)..(943)

; OTHER INFORMATION: n = A,T,C or G

US-09-833-381-1259

Query Match 72.0%; Score 18; DB 10; Length 943;  
Best Local Similarity 100.0%; Pred. No. 7.1;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 GTGCAGGACGTGACAAAT 25

Db 153 GTGCAGGACGTGACAAAT 136

##### RESULT 2

US-10-087-523-1

; Sequence 1, Application US/10087523

; Publication No. US20020197624A1

; GENERAL INFORMATION:

; APPLICANT: Klein, Robert D.

; APPLICANT: Brennan, Thomas J.

; TITLE OF INVENTION: METHODS OF CREATING CONSTRUCTS USEFUL FOR INTRODUCING

; SEQUENCES INTO EMBRYONIC STEM CELLS

us-09-787-562-1.1.rnpb

Mon Jan 6 15:20:17 2003

; FILE REFERENCE: 376472000200  
; CURRENT APPLICATION NUMBER: US/10/087,523  
; CURRENT FILING DATE: 2002-02-28  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/193,834  
; PRIOR FILING DATE: EARLIER FILING DATE: 1998-11-17  
; NUMBER OF SEQ ID NOS: 44  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 1  
; LENGTH: 4768  
; TYPE: DNA  
; ORGANISM: Plasmid vector  
US-10-087-523-1

Query Match 72.0%; Score 18; DB 9; Length 4768;  
Best Local Similarity 100.0%; Pred. No. 9;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
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DB 2824 GTGCAGGACGTGACAAAT 2841

RESULT 3  
US-09-816-790-1  
; Sequence 1, Application US/09816790  
; Patent No. US2002002225A1  
; GENERAL INFORMATION:  
; APPLICANT: Allen, Keith D.  
; APPLICANT: Phillips, Russell  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING  
; TITLE OF INVENTION: SULFOTRANSFERASE GENE DISRUPTIONS  
; FILE REFERENCE: R-855  
; CURRENT APPLICATION NUMBER: US/09/816,790  
; PRIOR FILING DATE: 2001-03-22  
; PRIOR APPLICATION NUMBER: US 60/191,240  
; PRIOR FILING DATE: 2000-03-22  
; PRIOR APPLICATION NUMBER: US 60/204,230  
; PRIOR FILING DATE: 2000-05-15  
; PRIOR APPLICATION NUMBER: US 60/223,173  
; PRIOR FILING DATE: 2000-08-07  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 4768  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Phage vector  
US-09-816-790-1

Query Match 72.0%; Score 18; DB 10; Length 4768;  
Best Local Similarity 100.0%; Pred. No. 9;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
|||||  
DB 2824 GTGCAGGACGTGACAAAT 2841

RESULT 4  
US-09-861-077-1  
; Sequence 1, Application US/09861077  
; Patent No. US20020023275A1  
; GENERAL INFORMATION:  
; APPLICANT: Leviten, Michael W.  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MATRIX  
; TITLE OF INVENTION: METALLOPROTEASE GENE DISRUPTIONS  
; FILE REFERENCE: R-15  
; CURRENT APPLICATION NUMBER: US/09/861,077  
; CURRENT FILING DATE: 2000-05-17  
; PRIOR APPLICATION NUMBER: US 60/204,972  
; PRIOR FILING DATE: 2000-05-17  
; PRIOR APPLICATION NUMBER: US 60/215,394

; PRIOR FILING DATE: 2000-06-29  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 4768  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Phage vector  
US-09-861-077-1

Query Match 72.0%; Score 18; DB 10; Length 4768;  
Best Local Similarity 100.0%; Pred. No. 9;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
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DB 2824 GTGCAGGACGTGACAAAT 2841

RESULT 5  
US-09-815-825-1  
; Sequence 1, Application US/09815825  
; Patent No. US2002002652A1  
; GENERAL INFORMATION:  
; APPLICANT: Allen, Keith D.  
; APPLICANT: Phillips, Russell  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING CGMP  
; TITLE OF INVENTION: PHOSPHODIESTERASE GENE DISRUPTIONS  
; FILE REFERENCE: R-849  
; CURRENT APPLICATION NUMBER: US/09/815,825  
; CURRENT FILING DATE: 2001-03-22  
; PRIOR FILING DATE: 2001-03-22  
; PRIOR APPLICATION NUMBER: US 60/191,142  
; PRIOR FILING DATE: 2000-03-22  
; PRIOR APPLICATION NUMBER: US 60/204,227  
; PRIOR FILING DATE: 2000-05-15  
; PRIOR APPLICATION NUMBER: US 60/216,765  
; PRIOR FILING DATE: 2000-07-06  
; PRIOR APPLICATION NUMBER: US 60/219,182  
; PRIOR FILING DATE: 2000-07-19  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 4768  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Phage vector  
US-09-815-825-1

Query Match 72.0%; Score 18; DB 10; Length 4768;  
Best Local Similarity 100.0%; Pred. No. 9;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 GTGCAGGACGTGACAAAT 25  
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DB 2824 GTGCAGGACGTGACAAAT 2841

RESULT 6  
US-09-815-935-1  
; Sequence 1, Application US/09815935  
; Patent No. US20020038466A1  
; GENERAL INFORMATION:  
; APPLICANT: Allen, Keith D.  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MAGNESIUM  
; TITLE OF INVENTION: DEPENDENT PROTEIN PHOSPHATASE GENE DISRUPTIONS  
; FILE REFERENCE: R-723  
; CURRENT APPLICATION NUMBER: US/09/815,935  
; CURRENT FILING DATE: 2001-03-22  
; PRIOR APPLICATION NUMBER: US 60/191,235  
; PRIOR FILING DATE: 2000-03-22  
; PRIOR APPLICATION NUMBER: US 60/216,249

; PRIOR FILING DATE: 2000-07-06  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 4768  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Phage vector  
US-09-815-935-1

Query Match 72.0%; Score 18; DB 10; Length 4768;  
Best Local Similarity 100.0%; Pred. No. 9;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 GTGCAGGACGTGACAAAT 25  
|||||  
Db 2824 GTGCAGGACGTGACAAAT 2841

RESULT 7  
US-09-815-944-1  
; Sequence 1, Application US/09815944  
; Patent No. US2002038467A1  
; GENERAL INFORMATION:  
; APPLICANT: Allen, Keith D.  
; APPLICANT: Matthews, William  
; APPLICANT: Moore, Mark  
; APPLICANT: Phillips, Russell  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MELANOCYTE  
; FILE REFERENCE: R-654  
; CURRENT APPLICATION NUMBER: US/09/815,944  
; CURRENT FILING DATE: 2001-03-22  
; PRIOR APPLICATION NUMBER: US 60/191,236  
; PRIOR FILING DATE: 2000-03-22  
; PRIOR APPLICATION NUMBER: US 60/215,214  
; PRIOR FILING DATE: 2000-06-29  
; PRIOR APPLICATION NUMBER: US 60/218,075  
; PRIOR FILING DATE: 2000-07-12  
; PRIOR APPLICATION NUMBER: US 60/219,167  
; PRIOR FILING DATE: 2000-07-19  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Phage vector  
US-09-815-944-1

Query Match 72.0%; Score 18; DB 10; Length 4768;  
Best Local Similarity 100.0%; Pred. No. 9;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 GTGCAGGACGTGACAAAT 25  
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Db 2824 GTGCAGGACGTGACAAAT 2841

RESULT 8  
US-10-087-523-2  
; Sequence 2, Application US/10087523  
; Publication No. US2002019762A1  
; GENERAL INFORMATION:  
; APPLICANT: Klein, Robert D.  
; APPLICANT: Brennan, Thomas J.  
; TITLE OF INVENTION: METHODS OF CREATING CONSTRUCTS USEFUL FOR INTRODUCING  
; FILE REFERENCE: 37647200200  
; CURRENT APPLICATION NUMBER: US/10/087,523  
; CURRENT FILING DATE: 2002-02-28

; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/193,834  
; PRIOR FILING DATE: EARLIER FILING DATE: 1998-11-17  
; NUMBER OF SEQ ID NOS: 44  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 2  
; LENGTH: 6355  
; TYPE: DNA  
; ORGANISM: Plasmid vector  
US-10-087-523-2

Query Match 72.0%; Score 18; DB 9; Length 6355;  
Best Local Similarity 100.0%; Pred. No. 9.4;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 GTGCAGGACGTGACAAAT 25  
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Db 4411 GTGCAGGACGTGACAAAT 4428

RESULT 9  
US-09-816-790-2  
; Sequence 2, Application US/09816790  
; Patent No. US2002022255A1  
; GENERAL INFORMATION:  
; APPLICANT: Allen, Keith D.  
; APPLICANT: Phillips, Russell  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING  
; FILE REFERENCE: R-855  
; CURRENT APPLICATION NUMBER: US/09/816,790  
; CURRENT FILING DATE: 2001-03-22  
; PRIOR APPLICATION NUMBER: US 60/191,240  
; PRIOR FILING DATE: 2000-03-22  
; PRIOR APPLICATION NUMBER: US 60/204,230  
; PRIOR FILING DATE: 2000-05-15  
; PRIOR APPLICATION NUMBER: US 60/223,173  
; PRIOR FILING DATE: 2000-08-07  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 2  
; LENGTH: 6355  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Phage vector  
US-09-816-790-2

Query Match 72.0%; Score 18; DB 10; Length 6355;  
Best Local Similarity 100.0%; Pred. No. 9.4;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 GTGCAGGACGTGACAAAT 25  
|||||  
Db 4411 GTGCAGGACGTGACAAAT 4428

RESULT 10  
US-09-861-077-2  
; Sequence 2, Application US/09861077  
; Patent No. US20020023275A1  
; GENERAL INFORMATION:  
; APPLICANT: Leviten, Michael W.  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MATRIX  
; FILE REFERENCE: R-15  
; CURRENT APPLICATION NUMBER: US/09/861,077  
; CURRENT FILING DATE: 2000-05-17  
; PRIOR APPLICATION NUMBER: US 60/204,972  
; PRIOR FILING DATE: 2000-05-17  
; PRIOR APPLICATION NUMBER: US 60/215,394  
; PRIOR FILING DATE: 2000-06-29  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 4.0

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; SEQ ID NO 2
; LENGTH: 6355
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-935-2

Query Match          72.0%; Score 18; DB 10; Length 6355;
Best Local Similarity 100.0%; Pred. No. 9.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db      4411 GTGCAGGACGTGACAAAT 4428

RESULT 11
US-09-815-825-2
; Sequence 2, Application US/09815825
; Patent No. US20020026652A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; APPLICANT: Phillips, Russell
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING cGMP
; TITLE OF INVENTION: PHOSPHODIESTERASE GENE DISRUPTIONS
; FILE REFERENCE: R-849
; CURRENT APPLICATION NUMBER: US/09/815,825
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,142
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/204,227
; PRIOR FILING DATE: 2000-05-15
; PRIOR APPLICATION NUMBER: US 60/216,765
; PRIOR FILING DATE: 2000-07-06
; PRIOR APPLICATION NUMBER: US 60/219,182
; PRIOR FILING DATE: 2000-07-19
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 6355
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-825-2

Query Match          72.0%; Score 18; DB 10; Length 6355;
Best Local Similarity 100.0%; Pred. No. 9.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      8 GTGCAGGACGTGACAAAT 25
        |||
Db      4411 GTGCAGGACGTGACAAAT 4428

RESULT 12
US-09-815-935-2
; Sequence 2, Application US/09815935
; Patent No. US20020038466A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MAGNESIUM
; TITLE OF INVENTION: DEPENDENT PROTEIN PHOSPHATASE GENE DISRUPTIONS
; FILE REFERENCE: R-723
; CURRENT APPLICATION NUMBER: US/09/815,935
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,235
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/216,249
; PRIOR FILING DATE: 2000-07-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 6355
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-935-2

Query Match          72.0%; Score 18; DB 10; Length 6355;
Best Local Similarity 100.0%; Pred. No. 9.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      8 GTGCAGGACGTGACAAAT 25
        |||
Db      4411 GTGCAGGACGTGACAAAT 4428

RESULT 13
US-09-815-944-2
; Sequence 2, Application US/09815944
; Patent No. US20020038467A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; APPLICANT: Matthews, William
; APPLICANT: Moore, Mark
; APPLICANT: Phillips, Russell
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MELANOCYTE
; TITLE OF INVENTION: STIMULATING HORMONE RECEPTOR GENE DISRUPTIONS
; FILE REFERENCE: R-654
; CURRENT APPLICATION NUMBER: US/09/815,944
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,236
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/215,214
; PRIOR FILING DATE: 2000-06-29
; PRIOR APPLICATION NUMBER: US 60/218,075
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/219,167
; PRIOR FILING DATE: 2000-07-19
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 6355
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-944-2

Query Match          72.0%; Score 18; DB 10; Length 6355;
Best Local Similarity 100.0%; Pred. No. 9.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      8 GTGCAGGACGTGACAAAT 25
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Db      4411 GTGCAGGACGTGACAAAT 4428

RESULT 14
US-09-987-601-1
; Sequence 1, Application US/09987601
; Patent No. US20020098223A1
; GENERAL INFORMATION:
; APPLICANT: MOULLIER, Philippe
; APPLICANT: DANOS, Olivier
; APPLICANT: HEARD, Jean-Michel
; APPLICANT: FERRY, Nicholas
; TITLE OF INVENTION: BIOCOMPATIBLE IMPLANT FOR THE EXPRESSION AND IN VIVO
; TITLE OF INVENTION: SECRETION OF A THERAPEUTIC SUBSTANCE
; FILE REFERENCE: 0660-0145-ODIV
; CURRENT APPLICATION NUMBER: US/09/987,601
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: 2001-11-15
; PRIOR APPLICATION NUMBER: 09/225,509
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; PRIOR FILING DATE: EARLIER FILING DATE: 1999-01-06
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/523,814
; PRIOR FILING DATE: EARLIER FILING DATE: 1996-01-19
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: FR 93/04700
; PRIOR FILING DATE: EARLIER FILING DATE: 1993-04-21
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: FR 93/09185
; PRIOR FILING DATE: EARLIER FILING DATE: 1993-07-26
; NUMBER OF SEQ ID NOS: 1
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 1
; LENGTH: 8388
; TYPE: DNA
; ORGANISM: mus musculus, Mo-MuLV, and other
US-09-987-601-1

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Query Match      72.0%; Score 18; DB 10; Length 8388;
Best Local Similarity 100.0%; Pred. No. 9.8;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      8 GTGCAGGACGTGACAAAT 25
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Db      2238 GTGCAGGACGTGACAAAT 2255

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# RESULT 15

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US-09-919-580-500
; Sequence 500, Application US/09919580
; Patent No. US20020110832A1
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Secrist, Heather
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
; TITLE OF INVENTION: AND DIAGNOSIS OF COLON CANCER
; FILE REFERENCE: 210121.552
; CURRENT APPLICATION NUMBER: US/09/919,580
; CURRENT FILING DATE: 2001-07-30
; NUMBER OF SEQ ID NOS: 934
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 500
; LENGTH: 279
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-919-580-500

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Query Match      68.0%; Score 17; DB 10; Length 279;
Best Local Similarity 80.0%; Pred. No. 18;
Matches 20; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

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QY      1 CGCGTCGGTGCAGGACGTGACAAAT 25
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Db      4 CGCGTCGACGACGAGGATGTAATAAAT 28

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Search completed: January 4, 2003, 01:06:05  
Job time : 12.4385 secs





GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:46 ; Search time 171.12 Seconds  
(without alignments)  
3231.380 Million cell updates/sec

Title: US-09-787-562-2

Perfect score: 19  
Sequence: 1 gtcggtcaggaagtgaca 19

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

GenEmbl.\*

1: gb.ba.\*

2: gb.htg.\*

3: gb.in.\*

4: gb.om.\*

5: gb.ov.\*

6: gb.pat.\*

7: gb.ph.\*

8: gb.pl.\*

9: gb.pr.\*

10: gb.ro.\*

11: gb.sts.\*

12: gb.sy.\*

13: gb.un.\*

14: gb.vi.\*

15: gb.ba.\*

16: em.fun.\*

17: em.hum.\*

18: em.in.\*

19: em.mu.\*

20: em.om.\*

21: em.or.\*

22: em.ov.\*

23: em.pat.\*

24: em.ph.\*

25: em.pl.\*

26: em.ro.\*

27: em.sts.\*

28: em.un.\*

29: em.vi.\*

30: em.htg\_hum.\*

31: em.htg\_inv.\*

32: em.htg\_other.\*

33: em.htg\_mus.\*

34: em.htg\_pln.\*

35: em.htg\_rod.\*

36: em.htg\_mam.\*

37: em.htg\_vrt.\*

38: em\_sy.\*

39: em.htgo\_hum.\*

40: em.htgo\_mus.\*

41: em.htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	% Match	Query Length	DB ID	Description
1	19	100.0	19	6	AX023660 Sequence
2	19	100.0	25	6	AX023659 Sequence
3	16.4	86.3	252715	2	AC127371 Mus muscu
4	16.4	86.3	338579	1	AP003004 Mesorhizo
5	16	84.2	1652	9	BC029804 Homo sapi
6	16	84.2	1777	6	AX463539 Sequence
7	16	84.2	50422	9	AL607085 Human DNA
8	16	84.2	118270	2	AL128395 Rattus no
9	16	84.2	121563	2	CNS07VQ4
10	16	84.2	200839	9	AC020763
11	15.8	83.2	461	6	AX439829
12	15.8	83.2	698	8	PPU80658
13	15.8	83.2	699	8	CRU80647
14	15.8	83.2	807	8	PPU80657
15	15.8	83.2	812	8	PPU80660
16	15.8	83.2	826	8	AF323739
17	15.8	83.2	827	8	PPU80656
18	15.8	83.2	828	8	PPU80653
19	15.8	83.2	828	8	PPU80661
20	15.8	83.2	869	8	AF287880
21	15.8	83.2	870	8	KSP406489
22	15.8	83.2	1052	8	AF357071
23	15.8	83.2	1228	9	HSU66579
24	15.8	83.2	1545	6	AX078628
25	15.8	83.2	1815	6	AR097359
26	15.8	83.2	19158	8	AF188714 Emericell
27	15.8	83.2	58439	2	AC100283
28	15.8	83.2	71596	2	AC100877
29	15.8	83.2	87022	2	AP001089
30	15.8	83.2	107336	2	AC128578
31	15.8	83.2	110000	2	AC009579_3
32	15.8	83.2	110000	2	AC124444_3
33	15.8	83.2	125515	2	AC108003
34	15.8	83.2	129302	2	AC083918
35	15.8	83.2	142362	2	AC036162
36	15.8	83.2	143239	2	AP005285
37	15.8	83.2	154926	9	AP005263
38	15.8	83.2	159272	2	AC131024
39	15.8	83.2	161674	2	AC015567
40	15.8	83.2	165591	9	AC019239
41	15.8	83.2	169124	2	AP001375
42	15.8	83.2	173368	9	AC091647
43	15.8	83.2	183030	2	AC130772
44	15.8	83.2	185479	2	AP001484
45	15.8	83.2	188050	1	AL646072

# ALIGNMENTS

RESULT 1	AX023660	Sequence 2	19 bp	DNA	linear	PAT 15-SEP-2000
AX023660	Sequence 2	from Patent WO0017371.				
LOCUS	AX023660					
DEFINITION	Sequence 2	from Patent WO0017371.				
ACCESSION	AX023660					
VERSION	AX023660.1	GI:10184021				
KEYWORDS						
SOURCE	Mus sp.					
ORGANISM	Mus sp.					
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;					
AUTHORS	1 (bases 1 to 19)					
TITLE	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.					
JOURNAL	Binley, K.M. and Naylor, S.					
	Polynucleotide constructs and uses thereof					
	Patent: WO 0017371-A 2 30-MAR-2000;					

BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD (GB)		McPherson, J.D. and Waterston, R.H.	
FEATURES		Direct Submission	
source		Submitted (17-JUL-2002) Genome Sequencing Center, 4444 Forest Park Parkway, St. Louis, MO 63108, USA	
BASE COUNT		----- Genome Center -----	
ORIGIN		Center: Washington University Genome Sequencing Center	
Query Match		Center code: WUGSC	
Best Local Similarity		Web site: http://genome.wustl.edu/gsc/index.shtml	
Matches		Contact: submission@wustl.wustl.edu	
19; Conservative		----- Project Information -----	
0; Mismatches		Center project name: M_BA0035D04	
0; Indels		----- Summary Statistics -----	
0; Gaps		Sequencing vector: M13; 0%	
0; Indels		Sequencing vector: plasmid; 100%	
0; Gaps		Chemistry: Dye-terminator ET; 0% of reads	
0; Indels		Chemistry: Dye-terminator Big Dye; 100% of reads	
0; Gaps		Assembly program: Phrap; version 0.990319	
0; Indels		Consensus quality: 246396 bases at least Q40	
0; Gaps		Consensus quality: 246893 bases at least Q30	
0; Indels		Consensus quality: 247291 bases at least Q20	
0; Gaps		Insert size: 218000; agarose-fp	
0; Indels		Insert size: 251715; sum-of-ctgigs	
0; Gaps		Quality coverage: 0.00 in Q20 bases; agarose-fp	
0; Indels		Quality coverage: 11.44 in Q20 bases; sum-of-ctgigs	
0; Gaps		* NOTE: This is a 'working draft' sequence. It currently	
0; Indels		* consists of 11 contigs. The true order of the pieces	
0; Gaps		* is not known and their order in this sequence record is	
0; Indels		* arbitrary. Gaps between the contigs are represented as	
0; Gaps		* runs of N, but the exact sizes of the gaps are unknown.	
0; Indels		* This record will be updated with the finished sequence	
0; Gaps		* as soon as it is available and the accession number will	
0; Indels		* be preserved.	
0; Gaps		* 1247: contig of 1247 bp in length	
0; Indels		* 1248: gap of unknown length	
0; Gaps		* 1347: contig of 2449 bp in length	
0; Indels		* 1348: gap of unknown length	
0; Gaps		* 3797: gap of unknown length	
0; Indels		* 3896: gap of unknown length	
0; Gaps		* 3897: contig of 3042 bp in length	
0; Indels		* 6938: gap of unknown length	
0; Gaps		* 7038: gap of unknown length	
0; Indels		* 7039: contig of 16260 bp in length	
0; Gaps		* 23298: gap of unknown length	
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0; Gaps		* 23399: contig of 16042 bp in length	
0; Indels		* 39440: gap of unknown length	
0; Gaps		* 39441: contig of 23707 bp in length	
0; Indels		* 39442: gap of unknown length	
0; Gaps		* 63247: gap of unknown length	
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0; Indels		* 108506: gap of unknown length	
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0; Gaps		* 139967: gap of unknown length	
0; Indels		* 140067: contig of 109286 bp in length	
0; Gaps		* 249352: gap of unknown length	
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Best Local Similarity	94.4%;	Pred. No. 2.2e+03;	
Matches	17;	Conservative 0;	Mismatches 1; Indels 0; Gaps 0;
OY	2	TCGGTCAGGACGTGACA 19	
Db	60050	TCGGTCAGGACGTGACA 60067	
RESULT 5			
LOCUS	BC029804	1652 bp	mRNA linear PRI 20-MAY-2002
DESCRIPTION	Homo sapiens, LOC146433, clone MGC:34647 IMAGE:5192904, mRNA.		
ACCESSION	BC029804	complete cds.	
VERSION	BC029804.1	GI:20987449	
KEYWORDS	MGC.		
SOURCE	Homo sapiens.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
TITLE	1 (bases 1 to 1652)		
JOURNAL	Strausberg,R.		
	Direct Submission		
	Submitted (06-MAY-2002), National Institutes of Health, Mammalian		
	Gene Collection (MGC), Cancer Genomics Office, National Cancer		
	Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,		
	USA		
REMARK	NIH-MGC project URL: http://mgc.nci.nih.gov		
COMMENT	Contact: MGC help desk		
	Email: qcaps-r@mail.nih.gov		
	Tissue Procurement: Life Technologies, Inc.		
	cDNA Library Preparation: Life Technologies, Inc.		
	cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)		
	DNA Sequencing by: Baylor College of Medicine Human Genome		
	Sequencing Center		
	Center code: BCM-HGSC		
	Web site: http://www.hgsc.bcm.tmc.edu/cdna/		
	Contact: amg@bcm.tmc.edu		
	Gunnaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Hale, S.M.,		
	Yoon, V.S., Kowis, C.R., Lawrence, S., Martin, R.G., Muzny, D.M.,		
	Richards, S., Gibbs, R.A.		
	Clone distribution: MGC clone distribution information can be found		
	through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov		
	Series: IRAK Plate: 50 Row: 1 Column: 18		
	This clone was selected for full length sequencing because it		
	passed the following selection criteria: Hexamer frequency ORF		
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CDS			

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SPSPPHSTGSRVPVRAQGEGLIP\*  
BASE COUNT 336 a 504 c 499 g 313 t  
ORIGIN

Query Match 84.2%; Score 16; DB 9; Length 1652;  
Best Local Similarity 100.0%; Pred. No. 5.2e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGACGACGCG 16  
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Db 692 GTCGGTGACGACGCG 707

RESULT 6  
AX463539  
LOCUS AX463539 1777 bp DNA linear PAT 15-JUL-2002  
DEFINITION Sequence 103 from Patent WO0248337.  
ACCESSION AX463539  
VERSION AX463539.1 GI:21886313  
KEYWORDS  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE  
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
TITLE Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.  
JOURNAL

SECRETED PROTEINS  
DUGGAN, B.M., YAO, M.G. and GRIFFIN, J.A.  
Patent: WO 0248337-A 103 20-JUN-2002;  
INCYTE GENOMICS INC (US)

FEATURES  
Location/Qualifiers  
source  
1..1777  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/notes="Incyte ID No: 7486536CB1"

BASE COUNT 331 a 555 c 548 g 343 t  
ORIGIN

Query Match 84.2%; Score 16; DB 6; Length 1777;  
Best Local Similarity 100.0%; Pred. No. 5.2e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGACGACGCG 16  
|||||  
Db 851 GTCGGTGACGACGCG 866

RESULT 7  
AL607085/c  
LOCUS AL607085 50422 bp DNA linear PRI 08-APR-2002  
DEFINITION Human DNA sequence from clone RP11-529L18 on chromosome 10,  
complete sequence.

ACCESSION AL607085  
VERSION AL607085.8 GI:20135753  
KEYWORDS HTG.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE  
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
TITLE Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.  
JOURNAL Heath, P.

COMMENT  
Submitted (08-APR-2002) Wellcome Trust Sanger Institute, Hinxton,  
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:  
humquery@sanger.ac.uk  
On Apr 10, 2002 this sequence version replaced gi:16974012.  
During sequence assembly data is compared from overlapping clones.  
Where differences are found these are annotated as variations  
together with a note of the overlapping clone name. Note that the  
variation annotation may not be found in the sequence submission  
corresponding to the overlapping clone, as we submit sequences with  
only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all  
regions were either double-stranded or sequenced with an alternate  
chemistry or covered by high quality data (i.e., phred quality >= 30);  
an attempt was made to resolve all sequencing problems, such  
as compressions and repeats; all regions were covered by at least  
one plasmid subclone or more than one M13 subclone; and the  
assembly was confirmed by restriction digest. The following  
abbreviations are used to associate primary accession numbers given  
in the feature table with their source databases: Em; EMBL; SW;  
SWISSPROT; Tr; TREMBL; Wp; WORMPEP; Information on the WORMPEP  
database can be found at  
http://www.sanger.ac.uk/projects/Celegans/wormpep This sequence  
was generated from part of bacterial clone contigs of human  
chromosome 10, constructed by the Sanger Centre Chromosome 10  
Mapping Group. Further information can be found at  
http://www.sanger.ac.uk/HGP/Chr10  
RP11-529L18 is from the library RPI1-11.2 constructed by the group  
of Pieter de Jong. For further details see  
http://www.chori.org/bacpac/home.htm  
VECTOR: pBACE3.6.

FEATURES  
Location/Qualifiers  
source  
1..50422  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/chromosome="10"  
/clone="RP11-529L18"  
/clone\_lib="RPI1-11.2"

BASE COUNT 12195 a 11534 c 12203 g 14490 t  
ORIGIN

Query Match 84.2%; Score 16; DB 9;  
Best Local Similarity 100.0%; Pred. No. 4e+03;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GGTGACGACGTGACA 19  
|||||  
Db 2520 GGTGACGACGTGACA 2505

RESULT 8  
AC128395/c  
LOCUS AC128395 118270 bp DNA linear HTG 19-JUL-2002  
DEFINITION Rattus norvegicus clone CH230-128B22, \*\*\* SEQUENCING IN PROGRESS  
\*\*\*, 43 unordered pieces.

ACCESSION AC128395  
VERSION AC128395.1 GI:21909054  
KEYWORDS HTG; HTGS\_PHASE1.  
SOURCE Rattus norvegicus.  
ORGANISM Rattus norvegicus

REFERENCE  
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
TITLE Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae;  
JOURNAL Rattus.

1 (bases 1 to 118270)  
Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-osman, F.R., Allen, C.,  
Alsbrooks, S.L., Amaratunge, H.C., Are, J.R., Ayele, M., Banks, T.,  
Barbaria, J., Benton, J., Bimarge, K., Blankenburg, K., Brown, D.,  
Bouck, J., Bowie, S., Brieva, M., Brown, E., Brown, M., Bryant, N.P.,  
Buhay, C., Burck, P., Burkett, C., Burrell, K.L., Byrd, N.C.,  
Carron, T.F., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D.,  
Chen, G., Chen, R., Chen, Z., Chowdhry, I., Christopoulos, C.,  
Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R.,  
Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A.,  
Delaney, K.R., Delgado, O., Denn, A.L., Ding, Y., Dinh, H.H.,  
Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J.,  
Earnhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Escotto, M.,  
Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P.,  
Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R.,  
Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K.,  
Harris, C., Harris, K., Hart, M., Havlak, P., Hawes, A., Hernandez, J.,  
Hernandez, O., Hodson, A., Hogues, M., Holloway, C., Hollins, B.,  
Homs, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Jackson, L.E.,  
Jacobson, B., Jia, X., Johnson, R., Jolivet, S., Joudah, S.,  
Karlsson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C.,

Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lewis, L. C., Lewis, L., Li, J., Li, Z., Lichte, O., Lue, C., Liu, J., Liu, W., Louise, H., Lozano, R. J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J., Maheshwari, M., Mapua, E., Martin, R., Martindale, A., Martinez, E., Massey, E., Mawhney, E., McLeod, M. P., Meador, M., Mei, G., Metzger, M., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Morgan, M., Morris, S., Moser, M., Neal, Z., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokenkwo, S., Oguh, M., Okwuonu, G., Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L. L., Quiles, M., Ren, Y., Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savary, G., Scherer, S., Scott, G., Shen, H., Shooshtari, N., Sisson, I., Sodergren, E., Sonaike, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Tabor, P., Tamerisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S., Usmani, K., Vasquez, L., Vera, V., Villalobos, D., Vinson, R., Wang, Q., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleczyk, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y. F., Zhou, J., Zorrilla, S., Nelson, D., Weinstein, G., and Gibbs, R.

Direct Submission  
Unpublished  
2 (bases 1 to 118270)  
Worley, K.C.  
Direct Submission  
Submitted (19-JUL-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

----- Genome Center  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: hgsc-help@bcm.tmc.edu

----- Project Information  
Center project name: GRZ1  
Center clone name: CH230-128B22  
----- Summary Statistics  
Sequencing vector: Plasmid  
Chemistry: Dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.990329  
Consensus quality: 69006 bases at least Q40  
Consensus quality: 71372 bases at least Q30  
Consensus quality: 73129 bases at least Q20  
-----

\* NOTE: Estimated insert size may differ from sequence length  
(see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 43 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1  
1020: contig of 1020 bp in length  
1021  
1120: gap of unknown length  
1121  
2392: contig of 1172 bp in length  
2393  
2392: gap of unknown length  
2393  
3673: contig of 1281 bp in length  
3674  
3773: gap of unknown length  
3774  
5076: contig of 1303 bp in length  
5077  
5176: gap of unknown length  
5177  
6770: contig of 1594 bp in length  
6771  
6870: gap of unknown length  
6871  
8638: contig of 1767 bp in length  
8639  
8737: gap of unknown length  
8738  
9864: contig of 1127 bp in length  
9865  
9964: gap of unknown length  
9965  
10986: contig of 1022 bp in length  
10987  
11086: gap of unknown length  
11087  
12691: contig of 1605 bp in length  
12692  
12791: gap of unknown length  
12792  
14343: contig of 1552 bp in length

14344  
14444  
14444: contig of 1278 bp in length  
15721  
15821: gap of unknown length  
15822  
17281: contig of 1460 bp in length  
17282  
17381: gap of unknown length  
17382  
18908: contig of 1527 bp in length  
18909  
19008: gap of unknown length  
19009  
20601: contig of 1593 bp in length  
20602  
20701: gap of unknown length  
20702  
22259: contig of 1558 bp in length  
22260  
22359: gap of unknown length  
22360  
23498: contig of 1139 bp in length  
23499  
23598: gap of unknown length  
23599  
25129: contig of 1531 bp in length  
25229: gap of unknown length  
25230  
26735: contig of 1506 bp in length  
26736  
26835: gap of unknown length  
26836  
28367: contig of 1532 bp in length  
28368  
28467: gap of unknown length  
28468  
29978: contig of 1511 bp in length  
29979  
30078: gap of unknown length  
30079  
31451: contig of 1373 bp in length  
31452  
33097: gap of unknown length  
33098  
33197: gap of unknown length  
33198  
34788: contig of 1591 bp in length  
34789  
34888: gap of unknown length  
36737: contig of 1849 bp in length  
36738  
36837: gap of unknown length  
36838  
38328: contig of 1491 bp in length  
38329  
38428: gap of unknown length  
38429  
40475: contig of 2047 bp in length  
40476  
40576: gap of unknown length  
40577  
42115: contig of 1540 bp in length  
42116  
42215: gap of unknown length  
42216  
43596: contig of 1381 bp in length  
43597  
43696: gap of unknown length  
43697  
45346: contig of 1650 bp in length  
45347  
45446: gap of unknown length  
45447  
47250: contig of 1804 bp in length  
47251  
47350: gap of unknown length  
47351  
49576: contig of 2226 bp in length  
49577  
49676: gap of unknown length  
49677  
52558: contig of 2882 bp in length  
52559  
52658: gap of unknown length  
52659  
54645: contig of 1987 bp in length  
54646  
54745: gap of unknown length  
54746  
58870: contig of 4125 bp in length  
58871  
58970: gap of unknown length  
58971  
63137: contig of 4167 bp in length  
63138  
63237: gap of unknown length  
63238  
68954: contig of 5717 bp in length  
68955  
69054: gap of unknown length  
69055  
74577: contig of 5523 bp in length  
74578  
74677: gap of unknown length  
74678  
81209: contig of 6532 bp in length  
81210  
81309: gap of unknown length  
81310  
86316: contig of 5007 bp in length  
86317  
91067: gap of unknown length  
91068  
91167: contig of 4651 bp in length  
91168  
10053: gap of unknown length  
10054  
100153: contig of 8886 bp in length  
100154  
107094: gap of unknown length  
107095  
107194: contig of 6941 bp in length  
107195  
118270: contig of 11076 bp in length.

Location/Qualifiers  
1. 118270  
/organism="Rattus norvegicus"  
/db\_xref="taxon:10116"  
/clone="CH230-128B22"

29719 a 22572 c 22749 g 29078 t 14152 others  
BASE COUNT  
ORIGIN

Query Match 84.2%; Score 16; DB 2; Length 118270;  
 Best Local Similarity 100.0%; Pred. No. 3.7e+03;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCGGTGCAGGACGTGA 17  
 |||||  
 Db 48602 TCGGTGCAGGACGTGA 48587

RESULT 9  
 CNS07YQ4/C  
 LOCUS  
 DEFINITION Oryza sativa chromosome 12 clone OJ1306\_H03, \*\*\* SEQUENCING IN  
 AL713904  
 ACCESSION  
 VERSION  
 KEYWORDS HTG: HTGS\_P2; HTGS\_ACTIVEFIN.  
 SOURCE  
 ORGANISM Oryza sativa

REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL

COMMENT  
 Submitted (06-MAY-2002) Genoscope - Centre National de Sequencage :  
 BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr  
 - Web : www.genoscope.cns.fr)  
 On May 8, 2002 this sequence version replaced gi:20160268.  
 IMPORTANT: This sequence is unfinished and does not necessarily  
 represent the correct sequence.  
 Work on the sequence is in progress and the release of this data is  
 based on the understanding that the sequence may change as work  
 continue. The sequence may be contaminated with foreign sequence  
 from E.coli, yeast, vector, phage, etc.  
 The nucleotide sequence of this BAC clone was generated by  
 combining Monsanto and Genoscope sequencing data.  
 \* NOTE: This is a 'working draft' sequence.  
 \* This sequence will be replaced  
 \* by the finished sequence as soon as it is available and  
 \* the accession number will be preserved.

FEATURES  
 source  
 Location/Qualifiers  
 1..121563  
 /organism="Oryza sativa"  
 /cultivar="Nipponbare"  
 /sub\_species="japonica"  
 /db\_xref="taxon:4530"  
 /chromosome="12"  
 /clone="OJ1306\_H03"  
 /clone.lib="Monsanto"

BASE COUNT 33858 a 26851 c 27291 g 33563 t  
 ORIGIN

Query Match 84.2%; Score 16; DB 2; Length 121563;  
 Best Local Similarity 100.0%; Pred. No. 3.7e+03;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTG 16  
 |||||  
 Db 71663 GTCGGTGCAGGACGTG 71648

RESULT 10  
 AC020763/C  
 LOCUS  
 DEFINITION Homo sapiens chromosome 16 clone RP11-394B2, complete sequence.

ACCESSION AC020763  
 VERSION AC020763.5 GI:22122869  
 KEYWORDS HTG.  
 SOURCE human.  
 ORGANISM Homo sapiens

REFERENCE  
 AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 TITLE Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.  
 JOURNAL 1 (bases 1 to 200839)  
 DOI Sequencing of Human Chromosome 16  
 REFERENCE Unpublished  
 2 (bases 1 to 200839)  
 TITLE DOE Joint Genome Institute.  
 JOURNAL Direct Submission  
 REFERENCE Submitted (09-JAN-2000) Production Sequencing Facility, DOE Joint  
 TITLE Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA  
 JOURNAL 3 (bases 1 to 200839)  
 REFERENCE DOE Joint Genome Institute.  
 TITLE Direct Submission  
 JOURNAL Submitted (06-AUG-2002) Production Sequencing Facility, DOE Joint  
 REFERENCE Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA  
 TITLE On Aug 6, 2002 this sequence version replaced gi:8576110.  
 JOURNAL Sequence Quality Assessment.

COMMENT  
 This entry has been annotated with sequence quality  
 estimates computed by the Phrap assembly program.  
 All manually edited bases have been reduced to quality zero.  
 Quality levels above 40 are expected to have less than  
 1 error in 10,000 bp.  
 Base-by-base quality values are not generally visible from the  
 GenBank flat file format but are available as part  
 of this entry's ASN.1 file.

-----  
 Sequence Quality Assessment:  
 This entry has been annotated with sequence quality  
 estimates computed by the Phrap assembly program.  
 All manually edited bases have been reduced to quality zero.  
 Quality levels above 40 are expected to have less than  
 1 error in 10,000 bp.  
 Base-by-base quality values are not generally visible from the  
 GenBank flat file format but are available as part  
 of this entry's ASN.1 file.

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 Location/Qualifiers  
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 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /chromosome="16"  
 /clone="RP11-394B2"  
 BASE COUNT 47780 a 54805 c 50908 g 47346 t  
 ORIGIN

Query Match 84.2%; Score 16; DB 9; Length 200839;  
 Best Local Similarity 100.0%; Pred. No. 3.6e+03;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTG 16  
 |||||  
 Db 116671 GTCGGTGCAGGACGTG 116656

RESULT 11  
 AX439829  
 LOCUS  
 DEFINITION Sequence 8244 from Patent WO0229113.  
 ACCESSION  
 VERSION AX439829  
 KEYWORDS AX439829.1 GI:21664640  
 SOURCE  
 ORGANISM Bacillus clausii.  
 REFERENCE 1  
 TITLE Bacillus clausii  
 Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.  
 Berka, R. and Clausen, I.G.  
 Methods for monitoring multiple gene expression

Mon Jan 6 15:20:22 2003

JOURNAL Patent: WO 0229113-A 8244 11-APR-2002;  
Novozymes Biotech, Inc. (US); Novozymes A/S (DK)  
FEATURES Location/Qualifiers  
source  
1..461  
/organism="Bacillus clausii"  
/db\_xref="taxon:79880"

BASE COUNT 124 a 94 c 118 g 125 t

ORIGIN

Query Match 83.2%; Score 15.8; DB 6; Length 461;  
Best Local Similarity 89.5%; Pred. No. 7.1e+03;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTGACA 19

Db 9 GTCGCTGCTGGACGTGACA 27

RESULT 12  
LOCUS PP080658 698 bp DNA linear PLN 05-JAN-1999  
DEFINITION Peniophora pini 25S nuclear ribosomal RNA gene, partial sequence.  
ACCESSION U80658  
VERSION U80658.1 GI:4098668  
KEYWORDS Peniophora pini.  
SOURCE Peniophora pini.  
ORGANISM Peniophora pini  
Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;  
Aphyllophorales; Lachnocladiaceae; Peniophora.

REFERENCE 1 (bases 1 to 698)  
Hallenberg, N. and Parmasto, E.  
Phylogenetic studies in species of Corticiaceae growing on branches  
of Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;  
Aphyllophorales; Lachnocladiaceae; Peniophora.

REFERENCE 2 (bases 1 to 698)  
Hallenberg, N.  
Direct Submission  
Submitted (02-DEC-1996) Systematic Botany, Botanical Institute,  
Carl Skottsbergs gata 22, Gothenburg S-413 19, Sweden  
Location/Qualifiers  
1..698  
/organism="Peniophora pini"  
/strain="FCUG 2399"  
/db\_xref="taxon:55353"  
1..698  
/product="25S ribosomal RNA"

FEATURES source

BASE COUNT 109 a 144 c 102 g 124 t

ORIGIN

Query Match 83.2%; Score 15.8; DB 8; Length 698;  
Best Local Similarity 89.5%; Pred. No. 6.9e+03;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTGACA 19

Db 56 GTCGCTGCTGGACGTGACA 74

RESULT 13  
LOCUS CR080647 699 bp DNA linear PLN 05-JAN-1999  
DEFINITION Corticium roseum 25S nuclear ribosomal RNA gene, partial sequence.  
ACCESSION U80647  
VERSION U80647.1 GI:4098657  
KEYWORDS Corticium roseum.  
SOURCE Corticium roseum.  
ORGANISM Corticium roseum  
Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;  
Aphyllophorales; Corticiaceae; Corticium.

REFERENCE 1 (bases 1 to 699)  
Hallenberg, N. and Parmasto, E.  
Phylogenetic studies in species of Corticiaceae growing on branches  
of Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;  
Aphyllophorales; Corticiaceae; Corticium.

REFERENCE 2 (bases 1 to 699)  
Hallenberg, N.  
Direct Submission  
Submitted (02-DEC-1996) Systematic Botany, Botanical Institute,  
Carl Skottsbergs gata 22, Gothenburg S-413 19, Sweden  
Location/Qualifiers  
1..699  
/organism="Corticium roseum"  
/strain="FCUG 2558"  
/db\_xref="taxon:55342"  
1..699  
/product="25S ribosomal RNA"

AUTHORS Hallenberg, N.  
TITLE Direct Submission  
JOURNAL Submitted (02-DEC-1996) Systematic Botany, Botanical Institute,  
Carl Skottsbergs gata 22, Gothenburg S-413 19, Sweden  
Location/Qualifiers  
1..699  
/organism="Corticium roseum"  
/strain="FCUG 2558"  
/db\_xref="taxon:55342"

FEATURES source

rRNA

BASE COUNT 113 a 140 c 97 g 130 t

ORIGIN

Query Match 83.2%; Score 15.8; DB 8; Length 699;  
Best Local Similarity 89.5%; Pred. No. 6.9e+03;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTGACA 19

Db 56 GTCGCTGCTGGACGTGACA 74

RESULT 14  
LOCUS PP080657 807 bp DNA linear PLN 05-JAN-1999  
DEFINITION Peniophora piceae 25S nuclear ribosomal RNA gene, partial sequence.  
ACCESSION U80657  
VERSION U80657.1 GI:4098667  
KEYWORDS Peniophora piceae.  
SOURCE Peniophora piceae.  
ORGANISM Peniophora piceae  
Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;  
Aphyllophorales; Lachnocladiaceae; Peniophora.

REFERENCE 1 (bases 1 to 807)  
Hallenberg, N. and Parmasto, E.  
Phylogenetic studies in species of Corticiaceae growing on branches  
of Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;  
Aphyllophorales; Lachnocladiaceae; Peniophora.

REFERENCE 2 (bases 1 to 807)  
Hallenberg, N.  
Direct Submission  
Submitted (02-DEC-1996) Systematic Botany, Botanical Institute,  
Carl Skottsbergs gata 22, Gothenburg S-413 19, Sweden  
Location/Qualifiers  
1..807  
/organism="Peniophora piceae"  
/strain="FCUG 2306"  
/db\_xref="taxon:55352"  
1..807  
/product="25S ribosomal RNA"

FEATURES source

rRNA

BASE COUNT 133 a 181 c 119 g 155 t

ORIGIN

Query Match 83.2%; Score 15.8; DB 8; Length 807;  
Best Local Similarity 89.5%; Pred. No. 6.8e+03;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTGACA 19

Db 165 GTCGCTGCTGGACGTGACA 183

RESULT 15  
LOCUS PP080660 812 bp DNA linear PLN 05-JAN-1999  
DEFINITION Peniophora proxima 25S nuclear ribosomal RNA gene, partial  
sequence.  
ACCESSION U80660  
VERSION U80660.1 GI:4098670  
KEYWORDS Peniophora proxima.  
SOURCE Peniophora proxima.  
ORGANISM Peniophora proxima  
Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;



REFERENCE Aphyllophorales; Lachnocladiaceae; Peniophora.  
 1. (bases 1 to 812)  
 AUTHORS Hallenberg, N. and Parmasto, E.  
 TITLE Phylogenetic studies in species of Corticiaceae growing on branches  
 JOURNAL Unpublished  
 REFERENCE 2 (bases 1 to 812)  
 AUTHORS Hallenberg, N.  
 TITLE Direct Submission  
 JOURNAL Submitted (02-DEC-1996) Systematic Botany, Botanical Institute,  
 Carl Skottsbergs gata 22, Gothenburg S-413 19, Sweden

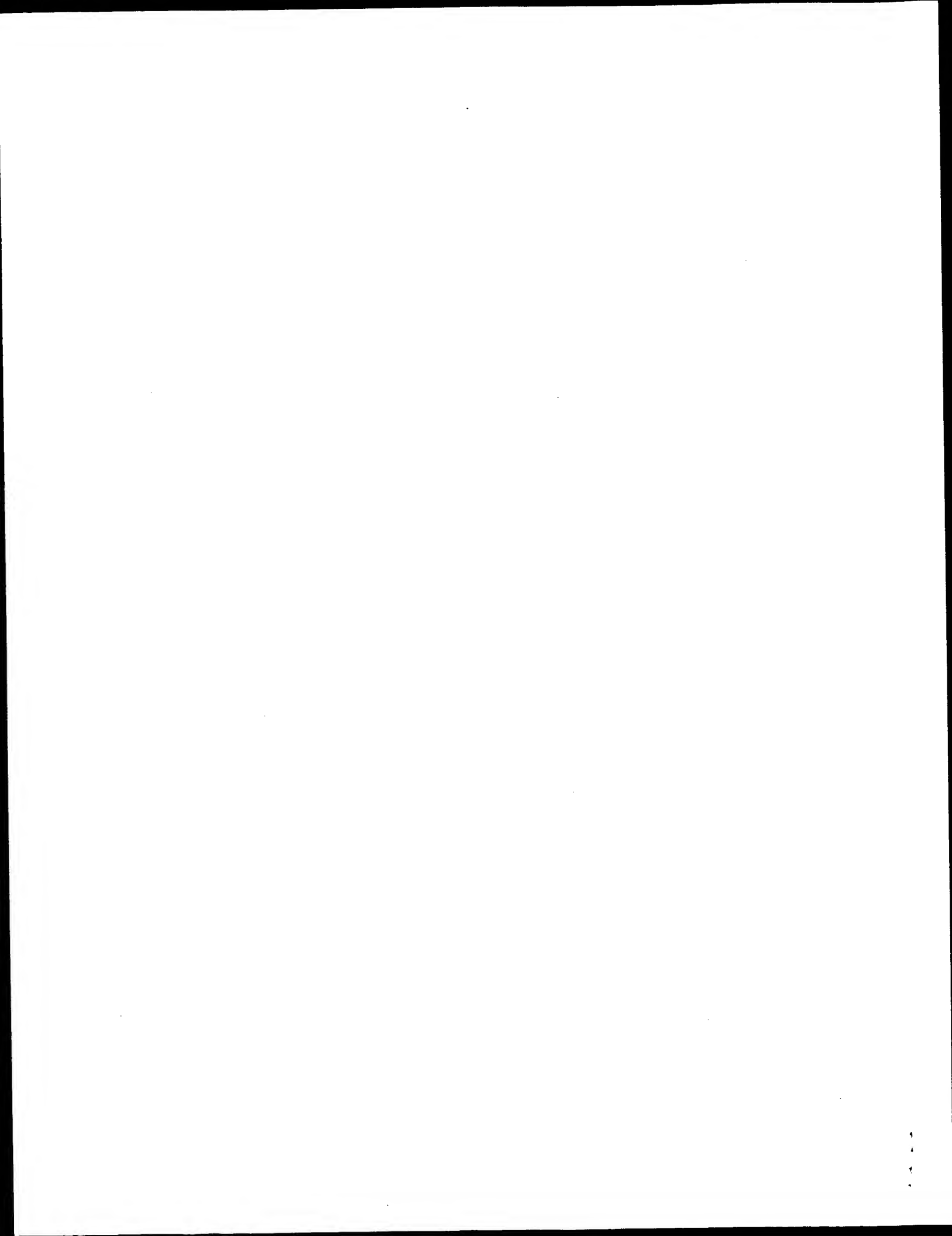
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 /strain="FCUG 1795"  
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BASE COUNT 135 a 181 c 119 g 158 t 219 others  
 ORIGIN

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 Best Local Similarity 89.5%; Pred. No. 6.8e+03;  
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GTCGGTGCAGGACGTGACA 19  
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 Db 171 GTCGGTGCAGGCGCGACA 189

Search completed: January 3, 2003, 23:54:35  
 Job time : 265.12 secs



GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:02 ; Search time 31.7666 Seconds  
(without alignments)  
1346.950 Million cell updates/sec

Title: us-09-787-562-2

Perfect score: 19  
Sequence: 1 gtcggtcaggacgtgaca 19

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	19	100.0	19	21	AAAA11994
2	19	100.0	25	21	AAAA11993
3	16	84.2	517	23	AAAS80329
4	16	84.2	633	23	AAAS80330
5	16	84.2	1624	21	AAC79886
6	15.8	83.2	461	24	ABK80953
7	15.8	83.2	1077	21	AAA30607
8	15.8	83.2	1077	21	AAA30720
9	15.8	83.2	1545	22	AAF61096

10	15.8	83.2	1815	20	AAZ06792
11	15.4	81.1	240	21	ABQ62698
12	15.4	81.1	1202	23	AAAS6562
13	15.4	81.1	1332	22	AAF60970
14	15.4	81.1	4403765	22	AAI99683
15	15.4	81.1	4411529	22	AAI99682
16	15	78.9	18	16	AAQ99459
17	15	78.9	18	21	AAA12053
18	15	78.9	18	21	AAA12054
19	15	78.9	19	22	AAF85326
20	15	78.9	21	22	AAH42134
21	15	78.9	24	16	AAQ99458
22	15	78.9	24	20	AAZ11422
23	15	78.9	24	21	AAA12007
24	15	78.9	24	22	AAC88980
25	15	78.9	43	22	AAH42138
26	15	78.9	72	20	AAZ11440
27	15	78.9	72	21	AAA12023
28	15	78.9	86	22	AAH42139
29	15	78.9	100	21	AAA12060
30	15	78.9	114	21	AAA12061
31	15	78.9	123	22	AAH42140
32	15	78.9	129	22	AAH42142
33	15	78.9	225	20	AAZ11399
34	15	78.9	225	21	AAA11997
35	15	78.9	229	20	AAZ11398
36	15	78.9	229	21	AAA11996
37	15	78.9	237	21	AAA12001
38	15	78.9	242	20	AAZ07789
39	15	78.9	242	21	AAA12016
40	15	78.9	243	20	AAZ11397
41	15	78.9	243	21	AAA11995
42	15	78.9	245	20	AAZ11438
43	15	78.9	249	21	AAA11999
44	15	78.9	269	20	AAZ11439
45	15	78.9			

#### ALIGNMENTS

RESULT 1  
AAAA11994  
ID AAA11994 standard; DNA; 19 BP.

AC AAA11994;

XX  
DT 14-AUG-2000 (first entry)

DE Murine PGK HRE truncated P18 DNA sequence.

XX HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;  
XX cardiant; cyclostatic; antiarthritic; gene therapy; ischaemia; arthritis;  
XX cardiovascular disease; peripheral arterial disease; cancer;  
XX phosphoglycerate kinase; PGK; murine; ds.  
OS Mus sp.  
XX  
XX  
XX WO200017371-A1.  
XX  
PD 30-MAR-2000.  
XX  
XX 22-SEP-1999; 99WO-GB03181.  
XX  
XX 23-SEP-1998; 98WO-CB02885.  
XX 28-JAN-1999; 99GB-0001906.  
XX 16-FEB-1999; 99GB-0003538.  
XX  
XX (OXFO-) OXFORD BIOMEDICA UK LTD.  
XX  
XX Binley KM, Naylor S;  
XX WPI; 2000-283595/24.

BCR673 (7 transmem  
Mycobacterium tube  
DNA encoding novel  
P. putida KT2440-a  
Mycobacterium tube  
Mycobacterium tube  
Hypoxia-inducible  
Murine PGK1 derive  
Murine promoter OB  
Nucleotide fragmen  
HRE element from t  
Hypoxia-inducible  
Murine HRE mPGK DN  
Murine hypoxic res  
Synapsin gene SIL  
HRE-containing enh  
Murine PGK HRE HIF  
HRE element contai  
Synapsin gene SIL  
ETAV U3 enhancer r  
ETAV U3 enhancer r  
Synapsin gene SIL  
PGK derived enhanc  
Murine PGK HRE der  
Murine PGK HRE der  
Promoter OBHrel us  
Murine PGK HRE der  
Murine PGK fragmen  
Murine PGK HRE der  
HIV derived synthe  
Murine PGK HRE der  
Synthetic promoter

hypoxia response element useful for driving expression of nucleic acids of interest in a cell under hypoxic conditions -

Disclosure; Page 11; 155pp; English.

This invention describes novel polynucleotide comprising at least 2 repeats of a hypoxia response element (HRE), where the hypoxia-inducible factor (HIF) consensus binding sites within each of the 2 repeats are separated by a spacer of at least 20 contiguous nucleotides. The products of the invention have vasotropic, cardiant, cytostatic and antiarthritic activity and can be used for gene therapy. The polynucleotides are useful for delivering nucleic acids of interest to mammalian cells. Lentiviral vectors are responsive to hypoxic agents and to agents that mimic hypoxia. This regulation can be harnessed in vitro to enhance the production of the vector and can be used in vivo to regulate gene expression in response to a physiological signal. The vectors have utility in disease, where ischaemia, including hypoxia, is a feature, e.g. cardiovascular disease, peripheral arterial disease, cancer and arthritis. The novel regulatory construct is capable of driving very high levels of transcription under conditions of hypoxia whilst providing only low basal levels of transcription under normal oxygen conditions. The polynucleotide construct targets cells within a tumor mass that are under conditions of hypoxia without affecting normal surrounding tissue. This sequence represents a murine phosphoglycerate kinase (PGK) HRE P24 DNA fragment as described in the method of the invention.

Sequence 25 BP; 6 A; 6 C; 9 G; 4 T; 0 other;

Query Match 100.0%; Score 19; DB 21; Length 25;  
Best Local Similarity 100.0%; Pred. No. 6.5; Mismatches 0; Indels 0; Gaps 0;  
Matches 19; Conservative 0;

QY 1 GTCGGTGCAGGACGTGACA 19  
|||||  
Db 4 GTCGGTGCAGGACGTGACA 22

RESULT 3  
AAS80329  
ID AAS80329 standard; cDNA; 517 BP.  
XX  
AC AAS80329;  
XX  
DT 13-FEB-2002 (first entry)  
XX  
DE DNA encoding novel human diagnostic protein #16133.  
XX  
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;  
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200175067-A2.  
XX  
PD 11-OCT-2001.  
XX  
PF 30-MAR-2001; 2001WO-US08631.  
XX  
PR 31-MAR-2000; 2000US-0540217.  
PR 23-AUG-2000; 2000US-0649167.  
XX  
PA (HYSE-) HYSEQ INC.  
XX  
PI Drmanac RT, Liu C, Tang YT;  
XX  
DR WPI; 2001-639362/73.  
XX  
PT P-PSDB; ABG16142.  
XX  
PT New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity -  
XX

Novel polynucleotide constructs comprising at least two repeats of a hypoxia response element useful for driving expression of nucleic acids of interest in a cell under hypoxic conditions -

Disclosure; Page 11; 155pp; English.

This invention describes novel polynucleotide comprising at least 2 repeats of a hypoxia response element (HRE), where the hypoxia-inducible factor (HIF) consensus binding sites within each of the 2 repeats are separated by a spacer of at least 20 contiguous nucleotides. The products of the invention have vasotropic, cardiant, cytostatic and antiarthritic activity and can be used for gene therapy. The polynucleotides are useful for delivering nucleic acids of interest to mammalian cells. Lentiviral vectors are responsive to hypoxic agents and to agents that mimic hypoxia. This regulation can be harnessed in vitro to enhance the production of the vector and can be used in vivo to regulate gene expression in response to a physiological signal. The vectors have utility in disease, where ischaemia, including hypoxia, is a feature, e.g. cardiovascular disease, peripheral arterial disease, cancer and arthritis. The novel regulatory construct is capable of driving very high levels of transcription under conditions of hypoxia whilst providing only low basal levels of transcription under normal oxygen conditions. The polynucleotide construct targets cells within a tumor mass that are under conditions of hypoxia without affecting normal surrounding tissue. This sequence represents a murine phosphoglycerate kinase (PGK) HRE truncated P18 DNA fragment as described in the method of the invention.

Sequence 19 BP; 4 A; 4 C; 8 G; 3 T; 0 other;

Query Match 100.0%; Score 19; DB 21; Length 19;  
Best Local Similarity 100.0%; Pred. No. 6.4; Mismatches 0; Indels 0; Gaps 0;  
Matches 19; Conservative 0;

QY 1 GTCGGTGCAGGACGTGACA 19  
|||||  
Db 1 GTCGGTGCAGGACGTGACA 19

RESULT 2  
AAAL1993  
ID AAAL1993 standard; DNA; 25 BP.  
XX  
AC AAAL1993;  
XX  
DT 14-AUG-2000 (first entry)  
XX  
DE Murine PGK HRE P42 DNA sequence.  
XX  
KW HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;  
KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;  
KW cardiovascular disease; peripheral arterial disease; cancer;  
KW phosphoglycerate kinase; PGK; murine; ds.  
XX  
OS Mus sp.  
XX  
PN WO200017371-A1.  
XX  
PD 30-MAR-2000.  
XX  
PF 22-SEP-1999; 99WO-GB03181.  
XX  
PR 23-SEP-1998; 98WO-GB02885.  
PR 28-JAN-1999; 99GB-0001906.  
PR 16-FEB-1999; 99GB-0003538.  
XX  
PA (OXFO-) OXFORD BIOMEDICA UK LTD.  
XX  
PI Binley KM, Naylor S;  
XX  
DR WPI; 2000-283595/24.  
XX  
PT Novel polynucleotide constructs comprising at least two repeats of a

PS Claim 1; SEQ ID No 16133; 103pp; English.  
 XX  
 CC The invention relates to isolated polynucleotide (I) and  
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
 CC and gene mapping, and in recombinant production of (II). The  
 CC polynucleotides are also used in diagnostics as expressed sequence tags  
 CC for identifying expressed genes. (I) is useful in gene therapy techniques  
 CC to restore normal activity of (II) or to treat disease states involving  
 CC (II). (II) is useful for generating antibodies against it, detecting or  
 CC quantitating a polypeptide in tissue, as molecular weight markers and as  
 CC a food supplement. (II) and its binding partners are useful in medical  
 CC imaging of sites expressing (II). (I) and (II) are useful for treating  
 CC disorders involving aberrant protein expression or biological activity.  
 CC The polypeptide and polynucleotide sequences have applications in  
 CC diagnostics, forensics, gene mapping, identification of mutations  
 CC responsible for genetic disorders or other traits to assess biodiversity  
 CC and to produce other types of data and products dependent on DNA and  
 CC amino acid sequences. AAS64197-AAS94564 represent novel human  
 CC diagnostic coding sequences of the invention.  
 CC Note: The sequence data for this patent did not appear in the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pct\_sequences.  
 XX  
 SQ Sequence 517 BP; 101 A; 137 C; 174 G; 105 T; 0 other;

Query Match 84.2%; Score 16; DB 23; Length 517;  
 Best Local Similarity 100.0%; Pred. No. 2.1e+02;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGGTGCAGGACGTG 16  
 |||||  
 Db 412 GTCGGTGCAGGACGTG 427

RESULT 4  
 AAS80330  
 ID AAS80330 standard; cDNA; 633 BP.  
 XX  
 AC AAS80330;  
 XX  
 DT 13-FEB-2002 (first entry)  
 XX  
 DE DNA encoding novel human diagnostic protein #16134.  
 XX  
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;  
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200175067-A2.  
 XX  
 PD 11-OCT-2001.  
 XX  
 PF 30-MAR-2001; 2001WO-US08631.  
 XX  
 PR 31-MAR-2000; 2000US-0540217.  
 XX  
 PR 23-AUG-2000; 2000US-0649167.  
 XX  
 PA (HYSE-) HYSEQ INC.  
 XX  
 PI Drmanac RT, Liu C, Tang YT;  
 XX  
 XX WPI; 2001-639362/73.  
 DR  
 DR P-PSDB; ABG16143.  
 XX  
 XX New isolated polynucleotide and encoded polypeptides, useful in  
 PT diagnostics, forensics, gene mapping, identification of mutations  
 PT responsible for genetic disorders or other traits and to assess  
 PT biodiversity -  
 XX  
 PS Claim 1; SEQ ID No 16134; 103pp; English.  
 XX

CC The invention relates to isolated polynucleotide (I) and  
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
 CC and gene mapping, and in recombinant production of (II). The  
 CC polynucleotides are also used in diagnostics as expressed sequence tags  
 CC for identifying expressed genes. (I) is useful in gene therapy techniques  
 CC to restore normal activity of (II) or to treat disease states involving  
 CC (II). (II) is useful for generating antibodies against it, detecting or  
 CC quantitating a polypeptide in tissue, as molecular weight markers and as  
 CC a food supplement. (II) and its binding partners are useful in medical  
 CC imaging of sites expressing (II). (I) and (II) are useful for treating  
 CC disorders involving aberrant protein expression or biological activity.  
 CC The polypeptide and polynucleotide sequences have applications in  
 CC diagnostics, forensics, gene mapping, identification of mutations  
 CC responsible for genetic disorders or other traits to assess biodiversity  
 CC and to produce other types of data and products dependent on DNA and  
 CC amino acid sequences. AAS64197-AAS94564 represent novel human  
 CC diagnostic coding sequences of the invention.  
 CC Note: The sequence data for this patent did not appear in the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pct\_sequences.  
 XX  
 SQ Sequence 633 BP; 147 A; 166 C; 189 G; 131 T; 0 other;

Query Match 84.2%; Score 16; DB 23; Length 633;  
 Best Local Similarity 100.0%; Pred. No. 2.1e+02;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGGTGCAGGACGTG 16  
 |||||  
 Db 447 GTCGGTGCAGGACGTG 462

RESULT 5  
 AAC79886  
 ID AAC79886 standard; cDNA; 1624 BP.  
 XX  
 AC AAC79886;  
 XX  
 DT 09-FEB-2001 (first entry)  
 XX  
 DE Human secreted protein encoding CDNA for gene 38.  
 XX  
 KW Human; secreted protein; cytostatic; antiarthritic; antiasthmatic;  
 KW immunosuppressive; antiarteriosclerotic; antiinflammatory; neurotropic;  
 KW neuroprotective; antidiabetic; tranquiliser; vulnary; antibacterial;  
 KW antipsoriatic; antiarrhythmic; antirheumatic; cardiant; anti-HIV;  
 KW autoimmune disorder; allergic condition; cardiovascular disorder;  
 KW cancer; neurological disease; tissue repair; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200055176-A2.  
 XX  
 PD 21-SEP-2000.  
 XX  
 PF 09-MAR-2000; 2000WO-US06057.  
 XX  
 PR 12-MAR-1999; 99US-0124142.  
 PR 11-JUN-1999; 99US-0138597.  
 PR 03-DEC-1999; 99US-0168666.  
 XX  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 XX  
 XX Rosen CA, Ruben SM, Komatsoulis G;  
 XX  
 XX WPI; 2000-638176/61.  
 DR  
 DR P-PSDB; AAB44867.  
 XX  
 XX Novel 49 human secreted proteins useful for diagnosis, prevention and  
 PT treatment of disorders including neurological, cell proliferative,  
 PT cardiovascular, and autoimmune/inflammatory disorders and microbial  
 PT infections -

Claim 1a; Page 355; 405pp; English.

This invention describes a novel isolated polypeptide (I) comprising an amino acid sequence at least 95 % identical to a polypeptide sequence selected from 49 polypeptides encoded by polynucleotide sequences included in American Type Culture Collection (ATCC) deposit number 203917, defined in the specification. The products of the invention have cytostatic, antiarthritic, antiasthmatic, immunosuppressive, neurotropic, antiarteriosclerotic, antiinflammatory, neuroprotective, antidiabetic, tranquiliser, vulnerary, antibacterial, antipsoriatic, antiarrhythmic, antineoplastic, anti-HIV activity. (I) or a nucleic acid (II) encoding (I) is useful for preventing, treating or ameliorating a medical condition and for diagnosing a pathological condition or susceptibility to the condition. (I) is useful for identifying a binding partner which affects the activity of the polypeptide and for identifying an active in a biological sample. (I), (II) or an antibody (IV) specific to (I) is also useful for treating or preventing a disease, disorder or condition associated with aberrant expression of (I). Diseases treated or diagnosed include immune disorders such as autoimmune diseases, blood protein disorders, anemia, allergic reactions and conditions such as asthma, organ rejection or graft-versus-host disease, inflammation, hyperproliferative disorders, cardiovascular disorders such as arterioarterial fistula, arrhythmias, arteriosclerosis, coronary thrombosis, organ regeneration, cancer, neovascular glaucoma, diabetic retinopathy, rheumatoid arthritis, psoriasis, diseases associated with increased apoptosis that include acquired immunodeficiency syndrome (AIDS), neurological diseases such as Parkinson's disease, viral, bacterial, fungal or parasitic diseases. They are also used to repair, replace or protect tissue damage by congenital defects, to treat trauma, in surgery, including cosmetic plastic surgery, to treat fibrosis, reperfusion injury or systemic cytokine damage, to stimulate chondrocyte growth, to prevent skin aging due to sunburn, to change a mammal's mental state or physical state by influencing biorhythms, cardiac rhythms, depression, memory, stress and for accelerating wound healing. (I), (II) and/or their agonist or antagonist are useful as food additives or preservatives to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamin, mineral or other nutritional components. (I) is useful for screening therapeutic compounds. (II) is useful in forensic biology for detecting DNA sequences and as diagnostic probes for detecting the presence of specific mRNA in a particular cell type.

Sequence 1624 BP; 303 A; 506 C; 503 G; 312 T; 0 other;

Query Match 84.2%; Score 16; DB 21; Length 1624;  
Best Local Similarity 100.0%; Pred No. 2.1e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY i GTCGGTGAGGACGCTG 16  
|||||  
Db 666 GTCGGTGAGGACGCTG 681

RESULT 6  
ABK80953  
ID ABK80953 standard; DNA; 461 BP.

XX AC ABK80953;  
XX  
DT 13-AUG-2002 (first entry)  
DE  
DE Bacillus clausii genomic sequence tag (GST) #3796.  
XX  
KW Differential gene expression; genomic sequenced tag; GST;  
KW altered culture condition; environmental stress;  
XX physiological provocation; ds.  
XX  
XX Bacillus clausii.  
XX OS  
XX WO200229113-A2.  
XX PN  
XX 11-APR-2002.  
XX DD

```

XX PF 12-OCT-1999; 99WO-US23938.
XX PR 13-OCT-1998; 98US-0170496.
XX PA (AREN-) ARENA PHARM INC.
XX PI Behan DP, Chalmers DT, Liaw CW;
XX DR WPI; 2000-329165/28.
XX DR P-PSDB; AAY90620.
XX PT Non-endogenous constitutively activated human G protein-coupled
XX PT receptors, useful for identifying agonists for use as pharmaceutical
XX PS agents.
XX PS Example 1; Page 131; 341pp; English.
XX CC The invention relates to constitutively active, non-endogenous versions
XX CC of endogenous human orphan G protein-coupled receptors (GPCRs, AAY90643-
XX CC AAY90677 and AAY90683-Y90687), and to DNA encoding them (AAA30709-A30743
XX CC and AAA30775-A30779). The mutant proteins of the invention contain a
XX CC mutation in a portion of the protein comprising intracellular loop 3
XX CC (IC3) and transmembrane domain 6 (TM6). A non-endogenous amino acid, X,
XX CC is substituted for an endogenous residue in TM6 to form a sequence
XX CC X-(AA)15-Pro. The endogenous proline in TM6 is selected from Lys, His, Arg
XX CC or Ala, and is preferably Lys. When the endogenous residue at this
XX CC position is Lys, this residue is replaced by His, Arg or preferably Ala.
XX CC The 15 amino acid stretch between the substituted amino acid and the Pro
XX CC may be endogenous, non-endogenous, or a mixture of endogenous and
XX CC non-endogenous residues. The constitutively active GPCRs are useful for
XX CC identifying antagonists, agonists and partial agonists for use as
XX CC pharmaceutical agents. The mutant proteins are also useful in research
XX CC settings for elucidating the roles of the receptors in normal and
XX CC diseased conditions. Antagonists for a particular GPCR are useful for
XX CC treating diseases and disorders associated with that receptor. Because
XX CC the novel mutant GPCRs are constitutively active, they can be used
XX CC directly for screening of compounds without the need for endogenous
XX CC ligands. The present sequence represents cDNA encoding a human wild-type
XX CC GPCR used in an exemplification of the invention. This was cloned and
XX CC subjected to site-directed mutagenesis (SDM) to generate DNA encoding
XX CC the corresponding mutant of the invention.
XX SQ Sequence 1077 BP; 142 A; 399 C; 320 G; 216 T; 0 other;

Query Match 83.2%; Score 15.8; DB 21; Length 1077;
Best Local Similarity 89.5%; Pred. No. 2.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTGACA 19
Db 549 GTCGGTGTGGCGGTGACA 567

RESULT 8
AAA30720
ID AAA30720 standard; DNA; 1077 BP.
AC AAA30720;
XX
XX 21-AUG-2000 (first entry)
XX
XX DNA encoding human mutant G protein-coupled receptor GPR20 (M240K).
XX G protein-coupled receptor; GPCR; constitutively active;
XX intracellular loop 3; transmembrane domain 6; drug screening;
XX agonist; antagonist; mutant; ss.
XX
XX Homo sapiens.
XX Synthetic.
XX WO200022129-A1.

```

```

XX PD 20-APR-2000.
XX PF 12-OCT-1999; 99WO-US23938.
XX PR 13-OCT-1998; 98US-0170496.
XX PA (AREN-) ARENA PHARM INC.
XX PI Behan DP, Chalmers DT, Liaw CW;
XX DR WPI; 2000-329165/28.
XX DR P-PSDB; AAY90654.
XX PT Non-endogenous constitutively activated human G protein-coupled
XX PT receptors, useful for identifying agonists for use as pharmaceutical
XX PS agents.
XX PS Example 2; Page 237; 341pp; English.
XX CC The invention relates to constitutively active, non-endogenous versions
XX CC of endogenous human orphan G protein-coupled receptors (GPCRs, AAY90643-
XX CC AAY90677 and AAY90683-Y90687), and to DNA encoding them (AAA30709-A30743
XX CC and AAA30775-A30779). The mutant proteins of the invention contain a
XX CC mutation in a portion of the protein comprising intracellular loop 3
XX CC (IC3) and transmembrane domain 6 (TM6). A non-endogenous amino acid, X,
XX CC is substituted for an endogenous residue in TM6 to form a sequence
XX CC X-(AA)15-Pro. The endogenous proline in TM6 is selected from Lys, His, Arg
XX CC or Ala, and is preferably Lys. When the endogenous residue at this
XX CC position is Lys, this residue is replaced by His, Arg or preferably Ala.
XX CC The 15 amino acid stretch between the substituted amino acid and the Pro
XX CC may be endogenous, non-endogenous, or a mixture of endogenous and
XX CC non-endogenous residues. The constitutively active GPCRs are useful for
XX CC identifying antagonists, agonists and partial agonists for use as
XX CC pharmaceutical agents. The mutant proteins are also useful in research
XX CC settings for elucidating the roles of the receptors in normal and
XX CC diseased conditions. Antagonists for a particular GPCR are useful for
XX CC treating diseases and disorders associated with that receptor. Because
XX CC the novel mutant GPCRs are constitutively active, they can be used
XX CC directly for screening of compounds without the need for endogenous
XX CC ligands. Sequences AAA30709- AAA30743 and AAA30775-A30779 represent DNAs
XX CC encoding the mutant human GPCRs of the invention.
XX SQ Sequence 1077 BP; 143 A; 399 C; 320 G; 215 T; 0 other;

Query Match 83.2%; Score 15.8; DB 21; Length 1077;
Best Local Similarity 89.5%; Pred. No. 2.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTGACA 19
Db 549 GTCGGTGTGGCGGTGACA 567

RESULT 9
AAF61096
ID AAF61096 standard; DNA; 1545 BP.
AC AAF61096;
XX
XX 16-MAY-2001 (first entry)
XX
XX P. putida KT2440-associated DNA ORF11200.
XX Transgenic plant; detection; probe; amplification; vaccine carrier;
XX microbial production strain; biological remediation; ds.
XX Pseudomonas putida.
XX DE19935088-A1.
XX 01-FEB-2001.

```

XX	27-JUL-1999;	99DE-1035088.	
XX	27-JUL-1999;	99DE-1035088.	
XX	(TIGR-) TIGR INST GENOMIC RES.		
XX	(QIUA-) QIAGEN GMBH.		
XX	(GBFB ) GES BIOTECHNOLOGISCHE FORSCHUNG MBH.		
XX	(DKFZ-) DKFZ DEUT KREBSFORSCHUNGSZENTRUM.		
XX	(MED1-) MEDIZINISCHE HOCHSCHULE HANNOVER.		
XX	WPI: 2001-192469/20.		
XX	New DNA sequences specific for <i>Pseudomonas putida</i> K72440, useful as		
XX	safe genetic engineering host, allow detection in presence of other		
XX	related bacteria -		
XX	Claim la; Page 149; 158pp; German.		
XX	This invention describes novel DNA sequences (1) for specific detection		
XX	of <i>Pseudomonas putida</i> K72440. The invention also describes (1)		
XX	recombinant expression vector containing (1); (2) prokaryotic or		
XX	eukaryotic cells transformed or transfected with (1) or the vector of		
XX	(1); (3) production of expression products by culturing cells of (2);		
XX	(4) expression products, or their fragments, of (1) and synthetic		
XX	proteins or peptides with the same sequences (A); (5) poly- or		
XX	mono-clonal antibodies (Ab) that react specifically with (A); (6)		
XX	hybridoma cells that produce the monoclonal Ab of (5); (7) transgenic		
XX	plants that contain transformed or transfected cells of (2); (8)		
XX	detecting K72440 using a labeled (1) or Ab as probe; and (9) DNA chips		
XX	carrying one or more (1). (1), and their fragments, are used as probes		
XX	to detect and isolate full-length cDNAs and/or to amplify such cDNAs by		
XX	polymerase chain reaction, and for production of transgenic plants. (1),		
XX	or antibodies that recognize their expression products, are used for		
XX	detecting the presence of K72440, particularly in presence of other,		
XX	even closely related, bacteria. K72440 is one of the bacteria classified		
XX	as safe, by the National Institutes of Health, for genetic engineering		
XX	work, e.g. as microbial production strains, for biological remediation		
XX	and as vaccine carriers. (1) are exclusive to K72440 with no significant		
XX	homology with sequences in other bacteria (specifically the closely		
XX	related pathogen <i>P. aeruginosa</i> ). Compared with other 'safe' bacteria, it		
XX	has greater catabolic activity and better survival in, and adaptation to,		
XX	the rhizosphere and soil.		
XX	Sequence 1545 BP; 255 A; 491 C; 496 G; 303 T; 0 other;		
XX	Query Match 83.2%; Score 15.8; DB 22; Length 1545;		
XX	Best Local Similarity 89.5%; Pred. No. 2.7e+02;		
XX	Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;		
QY	1 GTCGGTGCAGGACGTGACA 19		
Db	1494 GTCGGTGCAGGTGTGACA 1512		
RESULT 10			
AAZ06792			
ID	AAZ06792 standard; DNA; 1815 BP.		
AC			
XX	AAZ06792;		
XX			
XX	26-NOV-1999 (first entry)		
XX	ECR673 (7 transmembrane G-protein coupled receptor) gene.		
XX	7 transmembrane G-protein coupled receptor; HIV; viral infection; cancer;		
KW	Gilles de la Tourette's syndrome; neurological disorder; diabetes; asthma;		
KW	Parkinson's disease; Huntington's disease; signal transduction; ss.		
XX			
OS	Homo sapiens.		
XX			
XX	Key		
FH	Location/Qualifiers		



```

XX PF 16-APR-1999; 99WO-IB00740.
XX PR 16-APR-1998; 98US-0060756.
XX PA (INSP ) INST PASTEUR.
XX PI Cole S, Buchrieser-Brosch R, Gordon S, Billault A;
XX DR WPI; 2000-013262/01.
XX PT Isolation of polynucleotides from mycobacterial genomes, useful for
XX PT detection of Mycobacteria and for combating tuberculosis -
XX PS Claim 23; Page 62; 161pp; English.
XX CC The present invention describes a method for isolating a polynucleotide
XX CC of interest that is present or is expressed in a genome of a first
XX CC mycobacterium strain and that is absent or altered in a genome of a
XX CC second mycobacterium strain, which is different from the first strain
XX CC using a bacterial artificial chromosome (BAC) vector. Recombinant BAC
XX CC vectors, which are preferably immobilised, can be used to detect
XX CC mycobacterial nucleic acids (genomic DNA, cDNA or mRNA) in biological
XX CC samples. The polynucleotides identified are useful as probes or primers
XX CC for detecting a given mycobacterium of interest. By aligning the
XX CC polynucleotides contained in the recombinant BAC vectors it is possible
XX CC to physically map a polynucleotide of mycobacterial origin in a
XX CC biological sample. The methods and vectors from the present invention
XX CC are useful in providing information for combating tuberculosis. It is
XX CC possible to compare genomes between different strains or species and
XX CC their non-pathogenic strains or species counterparts. ABQ62492 to
XX CC ABQ63228 and AB881227 to AB881230 represent sequences used in the
XX CC exemplification of the present invention.
XX SQ Sequence 240 BP; 43 A; 70 C; 74 G; 53 T; 0 other;

Query Match 81.1%; Score 15.4; DB 21; Length 240;
Best Local Similarity 94.1%; Pred. No. 3.9e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 CGGTGCAGGACGTGACA 19
Db 110 CGGTGCAGGACGTGACA 94

RESULT 12
AA56562/c
ID AA56562 standard; cDNA; 1202 BP.
XX AC AA56562;
XX DT 13-FEB-2002 (first entry)
XX DE DNA encoding novel human diagnostic protein #2366.
XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX OS Homo sapiens.
XX PN WO200175067-A2.
XX PD 11-OCT-2001.
XX PF 30-MAR-2001; 2001WO-US08631.
XX PR 31-MAR-2000; 2000US-0540217.
XX PR 23-AUG-2000; 2000US-0649167.
XX PA (HYSE-) HYSEQ INC.
XX PI Drmanac RT, Liu C, Tang YT;
XX DR

WPI; 2001-639362/73.
P-PSDB; ABG02375.

New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity -

Claim 1; SEQ ID No 2366; 103pp; English.

The invention relates to isolated polynucleotide (I) and
polypeptide (II) sequences. (I) is useful as hybridisation probes,
polymerase chain reaction (PCR) primers, oligomers, and for chromosome
and gene mapping, and in recombinant production of (II). The
polynucleotides are also used in diagnostics as expressed sequence tags
for identifying expressed genes. (I) is useful in gene therapy techniques
to restore normal activity of (II) or to treat disease states involving
quantitating a polypeptide in tissue, as molecular weight markers and as
a food supplement. (II) and its binding partners are useful in medical
imaging of sites expressing (II). (I) and (II) are useful for treating
disorders involving aberrant protein expression or biological activity.
The polypeptide and polynucleotide sequences have applications in
diagnostics, forensics, gene mapping, identification of mutations
and to produce other types of data and products dependent on DNA and
amino acid sequences. AA564197-AA594564 represent novel human
diagnostic coding sequences of the invention.
Note: The sequence data for this patent did not appear in the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences.

Sequence 1202 BP; 290 A; 323 C; 304 G; 285 T; 0 other;

Query Match 81.1%; Score 15.4; DB 23; Length 1202;
Best Local Similarity 94.1%; Pred. No. 4.1e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCGGTGCAGGACGTGAC 18
Db 75 TGGGTGCAGGACGTGAC 59

RESULT 13
AAF60970
ID AAF60970 standard; DNA; 1332 BP.
XX AC AAF60970;
XX DT 16-MAY-2001 (first entry)
XX DE P. putida KT2440-associated DNA ORF00652.
XX KW Transgenic plant; detection; probe; amplification; vaccine carrier;
XX KW microbial production strain; biological remediation; ds.
XX OS Pseudomonas putida.
XX PN DE19935088-A1.
XX PD 01-FEB-2001.
XX PF 27-JUL-1999; 99DE-1035088.
XX PR 27-JUL-1999; 99DE-1035088.
XX PA (TIGR-) TIGR INST GENOMIC RES.
XX PA (QUIA-) QUIAGEN GMBH.
XX PA (GBFB ) GES BIOTECHNOLOGISCHE FORSCHUNG MBH.
XX PA (DKFZ-) DKFZ DEUT KREBSFORSCHUNGSZENTRUM.
XX PA (MEDI-) MEDIZINISCHE HOCHSCHULE HANNOVER.
XX DR WPI; 2001-192469/20.

```

XX New DNA sequences specific for *Pseudomonas putida* KT2440, useful as  
PT safe genetic engineering host, allow detection in presence of other  
PT related bacteria -  
XX  
PS Claim 1a; Page 17-18; 158pp; German.  
XX  
XX This invention describes novel DNA sequences (I) for specific detection  
CC of *Pseudomonas putida* KT2440. The invention also describes (1)  
CC recombinant expression vector containing (1); (2) prokaryotic or  
CC eukaryotic cells transformed or transfected with (1) or the vector of  
CC (1); (3) production of expression products by culturing cells of (2);  
CC (4) expression products, or their fragments, of (1) and synthetic  
CC proteins or peptides with the same sequences (A); (5) poly- or  
CC mono-clonal antibodies (Ab) that react specifically with (A); (6)  
CC hybridoma cells that produce the monoclonal Ab of (5); (7) transgenic  
CC plants that contain transformed or transfected cells of (2); (8)  
CC detecting KT2440 using a labeled (1) or Ab as probe; and (9) DNA chips  
CC carrying one or more (1), (I), and their fragments, are used as probes  
CC to detect and isolate full-length cDNAs and/or to amplify such cDNAs by  
CC polymerase chain reaction, and for production of transgenic plants, (I),  
CC or antibodies that recognize their expression products, are used for  
CC detecting the presence of KT2440, particularly in presence of other,  
CC even closely related, bacteria. KT2440 is one of the bacteria classified  
CC as safe, by the National Institutes of Health, for genetic engineering  
CC work, e.g. as microbial production strains, for biological remediation  
CC and as vaccine carriers. (I) are exclusive to KT2440 with no significant  
CC homology with sequences in other bacteria (specifically the closely  
CC related pathogen *P. aeruginosa*). Compared with other 'safe' bacteria, it  
CC has greater catabolic activity and better survival in, and adaptation to,  
CC the rhizosphere and soil.  
XX  
SQ Sequence 1332 BP; 182 A; 360 C; 486 G; 304 T; 0 other;  
  
Query Match 81.1%; Score 15.4; DB 22; Length 1332;  
Best Local Similarity 94.1%; Pred. No. 4.1e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
  
QY 1 GTCGGTGCAGGACGTGA 17  
Db 1230 GACCGTGCAGGACGTGA 1246  
|| |||||  
  
RESULT 14  
AAI99683  
ID AAI99683 standard; DNA; 4403765 BP.  
XX  
XX AAI99683;  
AC  
DT 15-JAN-2002 (first entry)  
XX  
DE Mycobacterium tuberculosis strain H37Rv genome SEQ ID NO 2.  
XX  
KW Mycobacterium tuberculosis; strain H37Rv; strain CDC 1551; genome;  
KW variation; epidemiology; patient treatment; epidemic monitoring; ds.  
XX  
OS Mycobacterium tuberculosis.  
XX  
XX US6294328-B1.  
XX  
PD 25-SEP-2001.  
XX  
PF 24-JUN-1998; 98US-0103840.  
XX  
PR 24-JUN-1998; 98US-0103840.  
XX  
XX (GENO-) INST GENOMIC RES.  
XX  
PI Fleischmann RD, White OR, Fraser CM, Venter JC;  
XX WPI; 2001-647261/74.  
XX  
XX Evaluating strain variation of Mycobacterium tuberculosis, comprises

PT determining the nucleotide sequence of the strain at positions in the  
PT genome corresponding to positions where *M. tuberculosis* strains CDC  
PT 1551 and H37Rv differ -  
XX  
PS Claim 4; SEQ ID NO 2; 3pp + Sequence Listing; English.  
XX  
XX The invention relates to evaluating strain variation within and between  
CC different populations of the tuberculosis bacterial pathogen,  
CC Mycobacterium tuberculosis or related Mycobacterium by determining the  
CC nucleotide sequence of the first strain at positions in the complete  
CC sequence of the genome that correspond to positions that differ in the  
CC nucleotide sequences of *M. tuberculosis* strains CDC 1551 (AAI99683) and  
CC H37Rv (AAI99682). The method is useful for evaluating strain variation of  
CC *M. tuberculosis* and has valuable application in the fields of  
CC tuberculosis genetics, epidemiology, patient treatment and epidemic  
CC monitoring  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from USPTO  
CC at [seqdata.uspto.gov/sequence.html?docID=6294328B1](http://seqdata.uspto.gov/sequence.html?docID=6294328B1).  
XX  
SQ Sequence 4403765 BP; 757105 A; 1447799 C; 1441301 G; 757371 T; 189 other;  
  
Query Match 81.1%; Score 15.4; DB 22; Length 4403765;  
Best Local Similarity 94.1%; Pred. No. 3.3e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
  
QY 1 GTCGGTGCAGGACGTGA 17  
Db 1664736 GTTGGTGCAGGACGTGA 1664752  
|| |||||  
  
RESULT 15  
AAI99682  
ID AAI99682 standard; DNA; 4411529 BP.  
XX  
XX AAI99682;  
AC  
XX 15-JAN-2002 (first entry)  
DT  
XX  
DE Mycobacterium tuberculosis strain H37Rv genome SEQ ID NO 1.  
XX  
KW Mycobacterium tuberculosis; strain H37Rv; strain CDC 1551; genome;  
KW variation; epidemiology; patient treatment; epidemic monitoring; ds.  
XX  
OS Mycobacterium tuberculosis.  
XX  
XX US6294328-B1.  
XX  
PD 25-SEP-2001.  
XX  
PF 24-JUN-1998; 98US-0103840.  
XX  
PR 24-JUN-1998; 98US-0103840.  
XX  
XX (GENO-) INST GENOMIC RES.  
XX  
PI Fleischmann RD, White OR, Fraser CM, Venter JC;  
XX WPI; 2001-647261/74.  
XX  
XX Evaluating strain variation of Mycobacterium tuberculosis, comprises  
PT determining the nucleotide sequence of the strain at positions in the  
PT genome corresponding to positions where *M. tuberculosis* strains CDC  
PT 1551 and H37Rv differ -  
XX  
PS Claim 3; SEQ ID NO 1; 3pp + Sequence Listing; English.  
XX  
XX The invention relates to evaluating strain variation within and between  
CC different populations of the tuberculosis bacterial pathogen,  
CC Mycobacterium tuberculosis or related Mycobacterium by determining the  
CC nucleotide sequence of the first strain at positions in the complete  
CC sequence of the genome that correspond to positions that differ in the  
CC nucleotide sequences of *M. tuberculosis* strains CDC 1551 (AAI99683) and  
CC nucleotide sequences of *M. tuberculosis* strains CDC 1551 (AAI99683) and

CC H37Rv (AAI99682). The method is useful for evaluating strain variation of  
CC M. tuberculosis and has valuable application in the fields of  
CC tuberculosis genetics, epidemiology, patient treatment and epidemic  
CC monitoring.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from USPTO  
CC at seqdata.uspto.gov/sequence.html?DocID=6294328B1.  
XX

SQ Sequence 4411529 BP; 758565 A; 1449983 C; 1444602 G; 758379 T; 0 other;

Query Match 81.1%; Score 15.4; DB 22; Length 4411529;  
Best Local Similarity 94.1%; Pred. No. 3.3e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GTCGGTGCAGGACGTGA 17  
|| |||||  
Db 1664602 GTTGGTGCAGGACGTGA 1664618

Search completed: January 3, 2003, 23:20:41  
Job time : 1049.77 secs



GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:17 ; Search time 252,634 Seconds  
(without alignments)  
1218.024 Million cell updates/sec

Title: US-09-787-562-2  
Perfect score: 19  
Sequence: 1 gtcggtgcaggacgtgaca 19

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :	EST :*	1:			
		em_estba.*	em_esthum.*	em_estin.*	em_estmu.*
		2: em_estba.*	2: em_esthum.*	3: em_estin.*	4: em_estmu.*
		3: em_estba.*	3: em_esthum.*	4: em_estin.*	5: em_estmu.*
		4: em_estba.*	4: em_esthum.*	5: em_estin.*	6: em_estmu.*
		5: em_estba.*	5: em_esthum.*	6: em_estin.*	7: em_estmu.*
		6: em_estba.*	6: em_esthum.*	7: em_estin.*	8: em_estmu.*
		7: em_estba.*	7: em_esthum.*	8: em_estin.*	9: gb_estl.*
		8: em_estba.*	8: em_esthum.*	9: gb_estl.*	10: gb_est2.*
		9: gb_estl.*	10: gb_est2.*	11: gb_htc.*	12: gb_est3.*
		10: gb_est2.*	11: gb_htc.*	12: gb_est3.*	13: gb_est4.*
		11: gb_htc.*	12: gb_est3.*	13: gb_est4.*	14: gb_est5.*
		12: gb_est3.*	13: gb_est4.*	14: gb_est5.*	15: em_estfun.*
		13: gb_est4.*	14: gb_est5.*	15: em_estfun.*	16: em_estom.*
		14: gb_est5.*	15: em_estfun.*	16: em_estom.*	17: gb_gss.*
		15: em_estfun.*	16: em_estom.*	17: gb_gss.*	18: em_gss_hum.*
		16: em_estom.*	17: gb_gss.*	18: em_gss_hum.*	19: em_gss_inv.*
		17: gb_gss.*	18: em_gss_hum.*	19: em_gss_inv.*	20: em_gss_pln.*
		18: em_gss_hum.*	19: em_gss_inv.*	20: em_gss_pln.*	21: em_gss_vrt.*
		19: em_gss_inv.*	20: em_gss_pln.*	21: em_gss_vrt.*	22: em_gss_fun.*
		20: em_gss_pln.*	21: em_gss_vrt.*	22: em_gss_fun.*	23: em_gss_mam.*
		21: em_gss_vrt.*	22: em_gss_fun.*	23: em_gss_mam.*	24: em_gss_mus.*
		22: em_gss_fun.*	23: em_gss_mam.*	24: em_gss_mus.*	25: em_gss_othr.*
		23: em_gss_mam.*	24: em_gss_mus.*	25: em_gss_othr.*	26: em_gss_pro.*
		24: em_gss_mus.*	25: em_gss_othr.*	26: em_gss_pro.*	27: em_gss_rod.*
		25: em_gss_othr.*	26: em_gss_pro.*	27: em_gss_rod.*	

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES			
Result No.	Score	Query Match Length DB ID	Description
c 1	16.4	86.3 567 13	BM140346 WHE0474_9
2	16.4	86.3 872 17	CNS03302
c 3	16.4	86.3 1021 17	AL225515 Tetraodon
c 4	16.4	86.3 1101 17	AL318185 Tetraodon
5	16	84.2 295 9	AL319040 Tetraodon
6	16	84.2 734 13	AI903141 QV-BT022- BI838592 603086272

7	16	84.2	786	13	BI819311
8	16	84.2	792	13	BI821807
9	16	84.2	799	13	BI916645
10	16	84.2	810	13	BI751993
11	16	84.2	819	13	BI829000
12	16	84.2	856	12	BG745622
c 13	16	84.2	884	9	AL537598
14	16	84.2	914	9	AL537599
15	16	84.2	1065	13	BM547868
16	16	84.2	1084	14	BM919043
17	16	84.2	1096	14	BM805234
18	15.8	83.2	392	10	AW436851
19	15.8	83.2	408	10	AW416022
20	15.8	83.2	442	17	AZ247056
c 21	15.8	83.2	448	10	BB839526
c 22	15.8	83.2	449	10	BB839535
23	15.8	83.2	475	12	BG334459
24	15.8	83.2	479	9	AA547770
25	15.8	83.2	531	17	AZ156676
26	15.8	83.2	556	10	AV616722
27	15.8	83.2	568	9	AJ273287
28	15.8	83.2	610	12	BG220636
29	15.8	83.2	784	17	BH400842
30	15.8	83.2	827	13	BI533157
31	15.8	83.2	907	17	CNS01KDC
c 32	15.4	81.1	250	10	BE011583
33	15.4	81.1	280	9	AJ493693
34	15.4	81.1	341	10	AW425923
c 35	15.4	81.1	405	9	AA933175
36	15.4	81.1	425	10	AV613228
37	15.4	81.1	442	10	BE481127
c 38	15.4	81.1	446	10	AW503758
c 39	15.4	81.1	453	10	AW520512
40	15.4	81.1	467	9	AL695680
c 41	15.4	81.1	469	9	AA998465
42	15.4	81.1	499	13	BI468593
43	15.4	81.1	507	13	BM588599
44	15.4	81.1	509	13	BM622212
45	15.4	81.1	519	13	BM610964

ALIGNMENTS

RESULT 1  
BM140346/c

LOCUS  
DEFINITION  
WHE0474\_g05\_n10zs Wheat Fusarium graminearum infected spike cDNA library Triticum aestivum cDNA clone WHE0474\_g05\_n10, mRNA sequence.

ACCESSION  
BM140346

VERSION  
BM140346.1

KEYWORDS  
EST.

SOURCE  
bread wheat.

ORGANISM  
Triticum aestivum

REFERENCE  
1 (bases 1 to 567)  
: Triticeae; Triticum.

AUTHORS  
Anderson, O.D., Chao, S., Han, P.S., Heinen, S., Hsia, C.C., Kang, Y., Kruger, W.M., Lazo, G.R., Miller, S., Muehlbauer, G.J., Miller, R., Pritsch, C., Rausch, C.J., Seaton, C.L., Tong, J.C., Vance, C. and Wilson, C.F.

TITLE  
The structure and function of the expressed portion of the wheat genomes - Fusarium graminearum infected spike cDNA library

JOURNAL  
COMMENT  
Contact: Olin Anderson  
US Department of Agriculture, Agriculture Research Service, Pacific West Area, Western Regional Research Center  
800 Buchanan Street, Albany, CA 94710, USA  
Tel: 5105595773  
Fax: 5105595818  
Email: oandersn@pw.usda.gov

567 bp mRNA linear EST 29-NOV-2001  
WHE0474\_g05\_n10zs Wheat Fusarium graminearum infected spike cDNA library Triticum aestivum cDNA clone WHE0474\_g05\_n10, mRNA sequence.

BI819311 603037760  
BI821807 603035886  
BI916645 603178586  
BI751993 603022105  
BI829000 603074878  
BG745622 602723868  
AL537598 AL537598  
AL537599 AL537599  
BM547868 AGENCOURT  
BM919043 AGENCOURT  
BM805234 AGENCOURT  
AW436851 77376 MAR  
AW416022 51001 MAR  
AZ247056 RPCI-23-4  
BB839526 BB839526  
BB839535 BB839535  
BG334459 PS28605.Y  
AA547770 EST188679  
AZ156676 SP\_0045.A  
AV616722 AV616722  
AJ273287 AJ273287  
BG220636 RST40423  
BH400842 AG-ND-171  
BI533157 603197755  
AL148193 Anopheles  
BE011583 CM4-BN022  
AJ493693 AJ493693  
AW425923 59015 MAR  
AA933175 UI-R-E0-d  
AV613228 AV613228  
BE481127 166443 BA  
AW503758 UI-HF-BN0  
AW520512 UI-R-BJ0p  
AL695680 AL695680  
AA998465 UI-R-C0-1  
BI468593 id89h12.x  
BM588599 170006873  
BM622212 170006874  
BM610964 170006591



```

BASE COUNT      309 a      217 c      229 g      255 t      11 others
ORIGIN

Query Match      86.3%; Score 16.4; DB 17; Length 1021;
Best Local Similarity 94.4%; Pred. No. 3.3e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GTCGGTCGACGACGTGAC 18
||| ||||| ||||| ||||| |||||
Db 127 GTTGGTCGACGACGTGAC 110

RESULT 4
CNS0535Z/c
LOCUS
DEFINITION      CNS0535Z 1101 bp DNA linear GSS 26-JUL-2000
Tetraodon nigroviridis genome survey sequence T3 end of clone
024D18 of library A from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION      AL319040
VERSION      AL319040.1 GI:9551924
KEYWORDS      GSS; genome survey sequence.
SOURCE      Tetraodon nigroviridis.
ORGANISM      Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Telostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontidae; Tetraodon.
REFERENCE      1 (bases 1 to 1101)
AUTHORS      Roest Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Saurin,W., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,A. and Weissenbach,J.
TITLE      Estimate of human gene number provided by genome-wide analysis
using Tetraodon nigroviridis DNA sequence
JOURNAL      Nat. Genet. 25 (2), 235-238 (2000)
MEDLINE      20296633
PUBMED      10835645
REFERENCE      2 (bases 1 to 1101)
AUTHORS      Crolius,H.R., Jaillon,O., Dasilva,C., Ozouf-Costaz,C., Fizames,C.,
Fischer,C., Bouneau,L., Billault,A., Quetier,F., Saurin,W.,
Brottier,P. and Weissenbach,J.
TITLE      Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL      Genome Res. 10 (7), 939-949 (2000)
MEDLINE      20359837
PUBMED      10899143
REFERENCE      3 (bases 1 to 1101)
AUTHORS      GenomeScope.
TITLE      Direct Submission
JOURNAL      Submitted (12-APR-2000)
COMMENT      This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
Location/Qualifiers
1..1101
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="024D18"
/clone_lib="A"
/note="Genoscope sequence ID : C0AA024DB09A1-end : T3"

BASE COUNT      277 a      268 c      308 g      227 t      21 others
ORIGIN

Query Match      86.3%; Score 16.4; DB 17; Length 1101;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GTCGGTCGACGACGTGAC 18
||| ||||| ||||| ||||| |||||
Db 94 GTCGGTCGACGACGTGAC 77

RESULT 5
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 734)

```

AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.  
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
 JOURNAL Unpublished (1999)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: [cgapbs-r@mail.nih.gov](mailto:cgapbs-r@mail.nih.gov)

Tissue Procurement: Life Technologies, Inc.  
 cDNA Library Preparation: Life Technologies, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>

Plate: LLAM1566 row: p column: 17  
 High quality sequence start: 5  
 High quality sequence stop: 728.

FEATURES

source

1. .734

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="IMAGE:522534"

/clone\_lib="NIH\_MGC\_120"

/lab\_host="DH10B"

/note="Organ: pooled pancreas and spleen; Vector: pCMV-SPORT6; Site.1: NotI; Site.2: EcoRV (destroyed); RNA source anonymous pool of spleen and pancreas from 28 yo male. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.5 kb, insert size range 1-2.5 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 025. Note: this is a NIH\_MGC Library."

BASE COUNT 144 a 210 c 231 g 149 t

ORIGIN

Query Match 84.2%; Score 16; DB 13; Length 734;

Best Local Similarity 100.0%; Pred. No. 4.6e-03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTCGAGCAGCTG 16

|||||

Db 412 GTCGGTCGAGCAGCTG 427

RESULT 7

BI819311

LOCUS

DEFINITION 603037760F1 NIH\_MGC\_115 Homo sapiens cDNA clone IMAGE:5178687 5', mRNA sequence.

ACCESSION BI819311

VERSION BI819311

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 786)

AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished (1999)

COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: [cgapbs-r@mail.nih.gov](mailto:cgapbs-r@mail.nih.gov)  
 Tissue Procurement: Life Technologies, Inc.  
 cDNA Library Preparation: Life Technologies, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>

Plate: LLAM1445 row: h column: 16

High quality sequence stop: 776.

FEATURES

source

1. 786

/organism="Homo sapiens"

/db\_xref="taxon:9606"  
 /clone="IMAGE:5178687"  
 /clone\_lib="NIH\_MGC\_115"  
 /lab\_host="DH10B"  
 /note="Organ: pooled brain, lung, testis; Vector: pCMV-SPORT6; Site.1: NotI; Site.2: EcoRV (destroyed); RNA source anonymous pool of 6 male brains, age range 23-27; 1 male lung, age 27; and 1 male testis, age 69. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.8 kb, insert size range 1-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 021. Note: this is a NIH\_MGC Library."

BASE COUNT 147 a 249 c 244 g 146 t

ORIGIN

Query Match 84.2%; Score 16; DB 13; Length 786;

Best Local Similarity 100.0%; Pred. No. 4.7e-03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTCGAGCAGCTG 16

|||||

Db 658 GTCGGTCGAGCAGCTG 673

RESULT 8

BI821807

LOCUS

DEFINITION 603035886F1 NIH\_MGC\_115 Homo sapiens cDNA clone IMAGE:5178686 5', mRNA sequence.

ACCESSION BI821807

VERSION BI821807

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 792)

AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished (1999)

COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: [cgapbs-r@mail.nih.gov](mailto:cgapbs-r@mail.nih.gov)  
 Tissue Procurement: Life Technologies, Inc.  
 cDNA Library Preparation: Life Technologies, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>

Plate: LLAM11440 row: 1 column: 21

High quality sequence stop: 789.

FEATURES

source

1. 792

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="IMAGE:5178686"

/clone\_lib="NIH\_MGC\_115"

/lab\_host="DH10B"

/note="Organ: pooled brain, lung, testis; Vector: pCMV-SPORT6; Site.1: NotI; Site.2: EcoRV (destroyed); RNA source anonymous pool of 6 male brains, age range 23-27; 1 male lung, age 27; and 1 male testis, age 69. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.8 kb, insert size range 1-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 021. Note: this is a NIH\_MGC Library."

BASE COUNT 146 a 249 c 248 g 149 t

ORIGIN



```

Query Match      84.2%; Score 16; DB 13; Length 792;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGCTG 16
      |||||
Db 641 GTCGGTGCAGGACGCTG 656

RESULT 9
BI916645
LOCUS
DEFINITION
603179586F1 NIH_MGC_121 Homo sapiens cDNA clone IMAGE:5242893 5',
mRNA sequence.
ACCESSION
BI916645
VERSION
BI916645.1 GI:16180607
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 799)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM11482 row: i column: 22
High quality sequence stop: 796.
Location/Qualifiers
1..799
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5242893"
/lab_host="NIH_MGC_121"
/note="Organ: brain; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: EcoRV (destroyed); RNA source anonymous pool of 3
fetal brains, female age 20 weeks, female age 24 weeks,
and male age 26 weeks. Library is oligo-dT primed and
directionally cloned (EcoRV site is destroyed upon
cloning). Average insert size 1.7 kb, insert size range
0.7-3.5 kb. Library is normalized and enriched for
full-length clones and was constructed by C. Gruber
(invitrogen). Research Genetics tracking code 017. Note:
this is a NIH_MGC Library."
BASE COUNT 152 a 254 c 247 g 146 t
ORIGIN

Query Match      84.2%; Score 16; DB 13; Length 799;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGCTG 16
      |||||
Db 676 GTCGGTGCAGGACGCTG 691

RESULT 10
BI751993
LOCUS
DEFINITION
603022103F1 NIH_MGC_114 Homo sapiens cDNA clone IMAGE:5192904 5',
mRNA sequence.
ACCESSION
BI751993
VERSION
BI751993.1 GI:15743571
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 810)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM11482 row: i column: 01
High quality sequence stop: 783.
Location/Qualifiers
1..810
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5192904"
/lab_host="NIH_MGC_114"
/note="Organ: brain; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: EcoRV (destroyed); RNA source anonymous pool of 6
male brains, age range 23-27 yo. Library is oligo-dT
primed and directionally cloned (EcoRV site is destroyed
upon cloning). Average insert size 1.5 kb, insert size
range 1-3 kb. Library is normalized and enriched for
full-length clones and was constructed by C. Gruber
(invitrogen). Research Genetics tracking code 019. Note:
this is a NIH_MGC Library."
BASE COUNT 154 a 257 c 251 g 148 t
ORIGIN

Query Match      84.2%; Score 16; DB 13; Length 810;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGCTG 16
      |||||
Db 692 GTCGGTGCAGGACGCTG 707

RESULT 11
BI829000
LOCUS
DEFINITION
603074878F1 NIH_MGC_119 Homo sapiens cDNA clone IMAGE:5166859 5',
mRNA sequence.
ACCESSION
BI829000
VERSION
BI829000.1 GI:15940550
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 819)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM11414 row: k column: 20

```

Query Match 84.2%; Score 16; DB 12; Length 856;  
 Best Local Similarity 100.0%; Pred. No. 4.8e+03;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTG 16  
 |||  
 DB 68 GTCGGTGCAGGACGTG 83

RESULT 13  
 AL537598/c  
 LOCUS  
 DEFINITION LTI\_FL013\_FBrnl Homo sapiens cDNA clone CS0DF026YK11 3  
 prime, mRNA sequence.  
 ACCESSION AL537598  
 VERSION AL537598.1 GI:12801091  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
 1 (bases 1 to 884)  
 Li W.B., Gruber C., Jessee, J. and Polayes, D.  
 Full-length cDNA libraries and normalization  
 Unpublished (2001)  
 Contact: Genoscope  
 Genoscope - Centre National de Sequencage  
 BP 191 91006 EVRY cedex - France  
 Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.  
 Location/Qualifiers  
 1..884  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone\_lib="LTI\_FL013\_FBrnl"  
 /dev\_stage="pooled tissue from post conception fetuses (20  
 week, 24 week and 26 week)"  
 /lab\_host="DH10B"  
 /note="Organ: Fetal brain; Vector: pCMVSPORT 6; 1st strand  
 cDNA was primed with a NotI-oligo(dT) primer. Five prime  
 end enriched, double-stranded cDNA was digested with Not I  
 and cloned into the Not I and Eco RV sites of the  
 pCMVSPORT 6 vector. Library was constructed by Life  
 Technologies. Contact : Feng Liang Life Technologies, a  
 division of Invitrogen 9800 Medical Center Drive Rockville  
 , Maryland 20850, USA Fax : (1) 301 610 8371 Email :  
 filiang@lifetech.com URL :  
 http://fulllength.invitrogen.com"

BASE COUNT 185 a 210 c 260 g 211 t 18 others  
 ORIGIN

Query Match 84.2%; Score 16; DB 9; Length 884;  
 Best Local Similarity 100.0%; Pred. No. 4.8e+03;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGGTGCAGGACGTG 16  
 |||  
 Db 767 GTCGGTGCAGGACGTG 752

RESULT 14  
 AL537599  
 LOCUS  
 DEFINITION LTI\_FL013\_FBrnl Homo sapiens cDNA clone CS0DF026YK11 5  
 prime, mRNA sequence.  
 ACCESSION AL537599  
 VERSION AL537599.1 GI:12801092  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL  
 COMMENT

Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
 1 (bases 1 to 914)  
 Li, W.B., Gruber, C., Jessee, J. and Polayres, D.  
 Full-length cDNA libraries and normalization  
 Unpublished (2001)  
 Contact: Genoscope  
 Genoscope - Centre National de Sequencage  
 BP 191 91006 Evry cedex - France  
 Email: [segref@genoscope.cns.fr](mailto:segref@genoscope.cns.fr), Web : [www.genoscope.cns.fr](http://www.genoscope.cns.fr).

FEATURES  
 source

1. .914  
 Location/Qualifiers  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="CS0DF26YK11"  
 /dev\_stage="LTI\_FL013\_FBrn1"  
 /week\_24="week 24, week and 26 week"  
 /lab\_host="DH108"  
 /note="Organ: Fetal brain; Vector: pCMVSPORT 6; 1st strand  
 cDNA was primed with a NotI-oligo(dT) primer. Five prime  
 end enriched, double-stranded cDNA was digested with Not I  
 and cloned into the Not I and Eco RV sites of the  
 pCMVSPORT 6 vector. Library was constructed by Life  
 Technologies. Contact : Feng Liang Life Technologies, a  
 division of Invitrogen 9800 Medical Center Drive Rockville  
 , Maryland 20850, USA Fax : (1) 301 610 8371 Email :  
[liang@life.com](mailto:liang@life.com) URL :  
<http://fulllength.invitrogen.com>"

BASE COUNT 170 a 291 c 277 g 176 t  
 ORIGIN

Query Match 84.2%; Score 16; DB 9; Length 914;  
 Best Local Similarity 100.0%; Pred. No. 4.8e+03;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTCGGTGCAGGACGTG 16  
 |||||||||||||  
 Db 659 GTCGGTGCAGGACGTG 674

RESULT 15  
 BM547868  
 LOCUS  
 DEFINITION  
 AGENCOURT\_6507246 NIH\_MGC\_124 Homo sapiens cDNA clone IMAGE:5727974  
 5', mRNA sequence.  
 BM547868  
 VERSION  
 BM547868.1 GI:18781998  
 EST.  
 SOURCE  
 human.

ORGANISM  
 Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
 1 (bases 1 to 1065)  
 NIH-MGC <http://mgs.nci.nih.gov/>.  
 National Institutes of Health, Mammalian Gene Collection (MGC)  
 Unpublished (1999)  
 Contact: Robert Strausberg, Ph.D.  
 Email: [cgapbs-r@mail.nih.gov](mailto:cgapbs-r@mail.nih.gov)  
 Tissue Procurement: Invitrogen  
 cDNA Library Preparation: Life Technologies, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Agencourt Bioscience Corporation  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
 Plate: L1AM12722 row: 0 column: 15  
 High quality sequence stop: 647.  
 Location/Qualifiers  
 1. .1065  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:5727974"  
 /clone\_lib="NIH\_MGC\_124"

FEATURES  
 source

/tissue\_type="hippocampus"  
 /lab\_host="DH108"  
 /note="Organ: brain; Vector: pCMV-SPORT6; Site\_1: EcoRV  
 (destroyed); Site\_2: NotI; RNA source male hippocampus,  
 age 27. Library is oligo-dT primed and directionally  
 cloned (EcoRV site is destroyed upon cloning). Average  
 insert size 1.4 kb, insert size range 0.9-4 kb. Library is  
 normalized and enriched for full-length clones and was  
 constructed by C. Gruber (Invitrogen). Research Genetics  
 tracking code 012."  
 BASE COUNT 197 a 345 c 324 g 199 t  
 ORIGIN

Query Match 84.2%; Score 16; DB 13;  
 Best Local Similarity 100.0%; Pred. No. 5e+03;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTCGGTGCAGGACGTG 16  
 |||||||||||||  
 Db 594 GTCGGTGCAGGACGTG 609

Search completed: January 4, 2003, 01:04:11  
 Job time : 256.634 secs



GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:53:16 : Search time 6.41325 Seconds  
(without alignments)  
908.566 Million cell updates/sec

Title: US-09-787-562-2

Perfect score: 19

Sequence: 1 gtcgtgcaggacgtgaca 19

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 15338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents\_NA.\*

1: /cgn2\_6/ptodata/2/ina/5A.COMB.seq.\*

2: /cgn2\_6/ptodata/2/ina/5B.COMB.seq.\*

3: /cgn2\_6/ptodata/2/ina/6A.COMB.seq.\*

4: /cgn2\_6/ptodata/2/ina/6B.COMB.seq.\*

5: /cgn2\_6/ptodata/2/ina/PCTUS.COMB.seq.\*

6: /cgn2\_6/ptodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15.8	83.2	1815	3	US-09-041-545-1
2	15.8	83.2	1815	3	US-09-327-925-1
3	15.4	81.1	240	4	US-09-060-756-197
4	15.4	81.1	4403765	4	US-09-103-840A-2
5	15.4	81.1	4411529	4	US-09-103-840A-1
6	15	78.9	18	2	US-08-693-174-3
7	15	78.9	18	2	US-08-853-236-1
8	15	78.9	18	4	US-09-253-738-3
9	15	78.9	24	2	US-08-693-174-2
10	15	78.9	24	4	US-09-253-738-2
11	15	78.9	41	2	US-08-853-236-2
12	15	78.9	1110	4	US-09-253-738-4
13	15	78.9	5382	4	US-09-479-122-21
14	15	78.9	7617	3	US-08-646-538-34
15	15	78.9	7617	3	US-09-503-222-34
16	15	78.9	8387	2	US-08-532-814-1
17	15	78.9	8388	4	US-09-225-509-1
18	15	78.9	9737	4	US-09-479-122-22
19	15	78.9	9737	4	US-09-479-122-23
20	15	78.9	9737	4	US-09-479-122-28
21	15	78.9	9737	4	US-09-479-122-28
22	15	78.9	9871	4	US-09-479-122-24
23	15	78.9	10060	4	US-09-479-122-25
24	14.8	77.9	1332	4	US-09-134-001C-1374
25	14.8	77.9	6854	4	US-09-194-905-7
26	14.8	77.9	43950	4	US-09-735-934A-3
27	14.8	77.9	152331	3	US-09-128-155-16

28	14.8	77.9	176373	3	US-09-128-155-17
29	14.4	75.8	444	4	US-09-134-001C-343
30	14.4	75.8	733	4	US-09-392-184-15
31	14.4	75.8	826	4	US-08-853-774-2
32	14.4	75.8	826	4	US-08-853-774-3
33	14.4	75.8	826	4	US-08-853-774-4
34	14.4	75.8	1447	4	US-09-484-970B-121
35	14.4	75.8	3691	4	US-09-211-704A-3
36	14.2	74.7	34	1	US-08-357-538-2
37	14.2	74.7	34	1	US-08-357-538-3
38	14.2	74.7	34	1	US-08-476-651-2
39	14.2	74.7	34	1	US-08-476-651-3
40	14.2	74.7	34	5	PCT-US93-10051-2
41	14.2	74.7	34	5	PCT-US93-10051-3
42	14.2	74.7	280	4	US-09-060-756-421
43	14.2	74.7	308	4	US-09-172-108-42
44	14.2	74.7	351	4	US-09-060-756-484
45	14.2	74.7	420	4	US-09-060-756-430

## ALIGNMENTS

## RESULT 1

US-09-041-545-1  
; Sequence 1, Application US/09041545  
; Patent No. 6071719  
; GENERAL INFORMATION:  
; APPLICANT: SATHE, GANESH M.  
; APPLICANT: HALSEY, WENDY S.  
; APPLICANT: MAO, JOYCE YUE  
; TITLE OF INVENTION: ECR 673 : A 7-TRANSMEMBRANE  
; TITLE OF INVENTION: G-PROTEIN COUPLED RECEPTOR  
; NUMBER OF SEQUENCES: 2  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: RATNER & PRESTIA  
; STREET: P.O. BOX 980  
; CITY: VALLEY FORGE  
; STATE: PA  
; COUNTRY: USA  
; ZIP: 19482  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette  
; COMPUTER: IBM Compatible  
; OPERATING SYSTEM: DOS  
; SOFTWARE: FastSeq for Windows Version 2.0  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/041,545  
; FILING DATE: 11-MAR-1998  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER:  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: PRESTIA, PAUL F  
; REGISTRATION NUMBER: 23,031  
; REFERENCE/DOCKET NUMBER: GP-70414  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 610-407-0700  
; TELEFAX: 610-407-0701  
; TELEX: 846169  
; INFORMATION FOR SEQ ID NO: 1:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 1815 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: cdna  
US-09-041-545-1

Query Match 83.2%; Score 15.8; DB 3; Length 1815;  
Best Local Similarity 89.5%; Pred. No. 47;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

RESULT 4  
US-09-103-840A-2  
; Sequence 2, Application US/09103840A  
; Patent No. 6294328  
; GENERAL INFORMATION:  
; APPLICANT: FLEISCHMAN, Robert D.  
; APPLICANT: WHITE, Owen R.  
; APPLICANT: FRASER, Claire M.  
; APPLICANT: VENTER, John C.  
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM  
; TITLE OF INVENTION: TUBERCULOSIS  
; FILE REFERENCE: 24366-20007.00  
; CURRENT APPLICATION NUMBER: US/09/103,840A  
; CURRENT FILING DATE: 1998-06-24  
; NUMBER OF SEQ ID NOS: 2  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 2  
; LENGTH: 4403765  
; TYPE: DNA  
; ORGANISM: Mycobacterium tuberculosis  
; FEATURE:  
; OTHER INFORMATION: CDC 1551  
; OTHER INFORMATION: "n" bases at various positions throughout the sequence  
; OTHER INFORMATION: represent a, t, c or g  
US-09-103-840A-2  
  
Query Match 81.1%; Score 15.4; DB 4; Length 4403765;  
Best Local Similarity 94.1%; Pred. No. 51;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
  
QY 1 GTCGGTGCAGGACGTGA 17  
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Db 1664736 GTTGGTCAGGACGTGA 1664752  
  
RESULT 5  
US-09-103-840A-1  
; Sequence 1, Application US/09103840A  
; Patent No. 6294328  
; GENERAL INFORMATION:  
; APPLICANT: FLEISCHMAN, Robert D.  
; APPLICANT: WHITE, Owen R.  
; APPLICANT: FRASER, Claire M.  
; APPLICANT: VENTER, John C.  
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM  
; TITLE OF INVENTION: TUBERCULOSIS  
; FILE REFERENCE: 24366-20007.00  
; CURRENT APPLICATION NUMBER: US/09/103,840A  
; CURRENT FILING DATE: 1998-06-24  
; NUMBER OF SEQ ID NOS: 2  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 1  
; LENGTH: 4411529  
; TYPE: DNA  
; ORGANISM: Mycobacterium tuberculosis  
; OTHER INFORMATION: H37Rv  
US-09-103-840A-1  
  
Query Match 81.1%; Score 15.4; DB 4; Length 4411529;  
Best Local Similarity 94.1%; Pred. No. 51;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
  
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Db 1664602 GTTGGTCAGGACGTGA 1664618  
  
RESULT 6  
US-08-693-174-3  
; Sequence 3, Application US/08693174A  
; Patent No. 5942434  
; GENERAL INFORMATION:  
; APPLICANT: Ratcliffe, Peter John

QY 1 GTCGGTGCAGGACGTGACA 19  
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Db 1169 GTCGGTGCAGGACGTGACA 1187  
  
RESULT 2  
US-09-327-925-1  
; Sequence 1, Application US/09327925A  
; Patent No. 6096868  
; GENERAL INFORMATION:  
; APPLICANT: SATHE, GANESH M.  
; APPLICANT: HALSEY, WENDY S.  
; APPLICANT: MAO, JOYCE YUE  
; TITLE OF INVENTION: COUPLED RECEPTOR  
; TITLE OF INVENTION: A 7 TRANSMEMBRANE G-PROTEIN  
; FILE REFERENCE: GP-70414-1  
; CURRENT APPLICATION NUMBER: US/09/327,925A  
; CURRENT FILING DATE: 1999-06-08  
; EARLIER APPLICATION NUMBER: 09/041,545  
; EARLIER FILING DATE: 1998-03-11  
; NUMBER OF SEQ ID NOS: 2  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 1  
; LENGTH: 1815  
; TYPE: DNA  
; ORGANISM: HOMO SAPIENS  
; FEATURE:  
; NAME/KEY: UNSURE  
; LOCATION: (16)  
US-09-327-925-1  
  
Query Match 83.2%; Score 15.8; DB 3; Length 1815;  
Best Local Similarity 89.5%; Pred. No. 47;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
  
QY 1 GTCGGTGCAGGACGTGACA 19  
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Db 1169 GTCGGTGCAGGACGTGACA 1187  
  
RESULT 3  
US-09-060-756-197/c  
; Sequence 197, Application US/09060756  
; Patent No. 6183957  
; GENERAL INFORMATION:  
; APPLICANT: Cole, Stewart  
; APPLICANT: Buchrieser-Brosch, Roland  
; APPLICANT: Gordon, Stephen  
; APPLICANT: Billault, Alain  
; TITLE OF INVENTION: METHOD FOR ISOLATING A POLYNUCLEOTIDE OF INTEREST FROM  
; TITLE OF INVENTION: THE GENOME OF A MYCOBACTERIUM USING A BAC-BASED DNA  
; TITLE OF INVENTION: LIBRARY APPLICATION TO THE DETECTION OF MYCOBACTERIA  
; FILE REFERENCE: 3495-0169  
; CURRENT APPLICATION NUMBER: US/09/060,756  
; CURRENT FILING DATE: 1998-04-16  
; NUMBER OF SEQ ID NOS: 743  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 197  
; LENGTH: 240  
; TYPE: DNA  
; ORGANISM: Mycobacterium tuberculosis  
US-09-060-756-197  
  
Query Match 81.1%; Score 15.4; DB 4; Length 240;  
Best Local Similarity 94.1%; Pred. No. 68;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
  
QY 3 CGGTGCAGGACGTGACA 19  
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Db 110 CGGTGCAGGACGTGACA 94

; APPLICANT: Firth, John David  
; APPLICANT: Harris, Adrian Llewellyn  
; APPLICANT: Pugh, Christopher William  
; APPLICANT: Stratford, Ian James  
; TITLE OF INVENTION: Targeting Gene Therapy  
; FILE REFERENCE: 08/693174  
; CURRENT APPLICATION NUMBER: US/08/693,174A  
; CURRENT FILING DATE: 1996-12-12  
; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 3  
; LENGTH: 18  
; TYPE: DNA  
; ORGANISM: Murinae gen. sp.  
US-08-693-174-3

Query Match 78.9%; Score 15; DB 2; Length 18;  
Best Local Similarity 100.0%; Pred. No. 98;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19  
Db 4 GTGCAGGACGTGACA 18  
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RESULT 7  
US-08-853-236-1  
; Sequence 1, Application US/08853236  
; Patent No. 5952226  
; GENERAL INFORMATION:  
; APPLICANT: Aebischer, Patrick  
; APPLICANT: Deglon, Nicole  
; APPLICANT: Regulier, Etienne  
; APPLICANT: Rinsch, Christopher  
; TITLE OF INVENTION: DEVICE AND METHOD FOR DELIVERY OF ERYTHROPOIETIN  
; FILE REFERENCE: Modex 004 divisional PGK-1 HRE  
; CURRENT APPLICATION NUMBER: US/08/853,236  
; CURRENT FILING DATE: 1997-05-09  
; NUMBER OF SEQ ID NOS: 2  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 1  
; LENGTH: 18  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: PGK-1 hypoxia  
US-08-853-236-1

Query Match 78.9%; Score 15; DB 2; Length 18;  
Best Local Similarity 100.0%; Pred. No. 98;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19  
Db 4 GTGCAGGACGTGACA 18  
|||||

RESULT 8  
US-09-253-738-3  
; Sequence 3, Application US/09253738  
; Patent No. 6265390  
; GENERAL INFORMATION:  
; APPLICANT: Ratcliffe, Peter John  
; APPLICANT: Firth, John David  
; APPLICANT: Harris, Adrian Llewellyn  
; APPLICANT: Pugh, Christopher William  
; APPLICANT: Stratford, Ian James  
; TITLE OF INVENTION: Targeting Gene Therapy  
; FILE REFERENCE: 08/693174  
; CURRENT APPLICATION NUMBER: US/09/253,738  
; CURRENT FILING DATE: 1999-02-22  
; NUMBER OF SEQ ID NOS: 5

; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 3  
; LENGTH: 18  
; TYPE: DNA  
; ORGANISM: Murinae gen. sp.  
US-09-253-738-3

Query Match 78.9%; Score 15; DB 4; Length 18;  
Best Local Similarity 100.0%; Pred. No. 98;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19  
Db 4 GTGCAGGACGTGACA 18  
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RESULT 9  
US-08-693-174-2  
; Sequence 2, Application US/08693174A  
; Patent No. 5942434  
; GENERAL INFORMATION:  
; APPLICANT: Ratcliffe, Peter John  
; APPLICANT: Firth, John David  
; APPLICANT: Harris, Adrian Llewellyn  
; APPLICANT: Pugh, Christopher William  
; APPLICANT: Stratford, Ian James  
; TITLE OF INVENTION: Targeting Gene Therapy  
; FILE REFERENCE: 08/693174  
; CURRENT APPLICATION NUMBER: US/08/693,174A  
; CURRENT FILING DATE: 1996-12-12  
; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 2  
; LENGTH: 24  
; TYPE: DNA  
; ORGANISM: Murinae gen. sp.  
US-08-693-174-2

Query Match 78.9%; Score 15; DB 2; Length 24;  
Best Local Similarity 100.0%; Pred. No. 99;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19  
Db 7 GTGCAGGACGTGACA 21  
|||||

RESULT 10  
US-09-253-738-2  
; Sequence 2, Application US/09253738  
; Patent No. 6265390  
; GENERAL INFORMATION:  
; APPLICANT: Ratcliffe, Peter John  
; APPLICANT: Firth, John David  
; APPLICANT: Harris, Adrian Llewellyn  
; APPLICANT: Pugh, Christopher William  
; APPLICANT: Stratford, Ian James  
; TITLE OF INVENTION: Targeting Gene Therapy  
; FILE REFERENCE: 08/693174  
; CURRENT APPLICATION NUMBER: US/09/253,738  
; CURRENT FILING DATE: 1999-02-22  
; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 2  
; LENGTH: 24  
; TYPE: DNA  
; ORGANISM: Murinae gen. sp.  
US-09-253-738-2

Query Match 78.9%; Score 15; DB 4; Length 24;  
Best Local Similarity 100.0%; Pred. No. 99;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 5 GTGCAGGACGTGACA 19
Db 7 GTGCAGGACGTGACA 21

RESULT 11
US-08-853-236-2
; Sequence 2, Application US/08853236
; Patent No. 5952226
; GENERAL INFORMATION:
; APPLICANT: Aebischer, Patrick
; APPLICANT: Degion, Nicole
; APPLICANT: Regulier, Etienne
; APPLICANT: Rinsch, Christopher
; TITLE OF INVENTION: DEVICE AND METHOD FOR DELIVERY OF ERYTHROPOIETIN
; FILE REFERENCE: Modex 004 divisional PKG-1 HRE
; CURRENT APPLICATION NUMBER: US/08/853,236
; CURRENT FILING DATE: 1997-05-09
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2
; LENGTH: 41
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: PKG-1 hypoxia
; OTHER INFORMATION: responsive element tandem repeat
US-08-853-236-2

Query Match 78.9%; Score 15; DB 2; Length 41;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 27 GTGCAGGACGTGACA 41

RESULT 12
US-08-693-174-4
; Sequence 4, Application US/08693174A
; Patent No. 5942434
; GENERAL INFORMATION:
; APPLICANT: Ratcliffe, Peter John
; APPLICANT: Firth, John David
; APPLICANT: Harris, Adrian Llewellyn
; APPLICANT: Pugh, Christopher William
; APPLICANT: Stratford, Ian James
; TITLE OF INVENTION: Targeting Gene Therapy
; FILE REFERENCE: 08/693174
; CURRENT APPLICATION NUMBER: US/08/693,174A
; CURRENT FILING DATE: 1996-12-12
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 1110
; TYPE: DNA
; ORGANISM: Murinae gen. sp.
US-08-693-174-4

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Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 637 GTGCAGGACGTGACA 651

RESULT 13
US-09-253-738-4
; Sequence 4, Application US/09253738
; Patent No. 6265390
; GENERAL INFORMATION:
; APPLICANT: Ratcliffe, Peter John
; APPLICANT: Firth, John David
; APPLICANT: Harris, Adrian Llewellyn
; APPLICANT: Pugh, Christopher William
; APPLICANT: Stratford, Ian James
; TITLE OF INVENTION: Targeting Gene Therapy
; FILE REFERENCE: 08/693174
; CURRENT APPLICATION NUMBER: US/09/253,738
; CURRENT FILING DATE: 1999-02-22
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 1110
; TYPE: DNA
; ORGANISM: Murinae gen. sp.
US-09-253-738-4

Query Match 78.9%; Score 15; DB 4; Length 5382;
Best Local Similarity 100.0%; Pred. No. 1.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 899 GTGCAGGACGTGACA 913

RESULT 14
US-09-479-122-21
; Sequence 21, Application US/09479122
; Patent No. 6410266
; GENERAL INFORMATION:
; APPLICANT: HARRINGTON, JOHN J.
; APPLICANT: SHEREF, BRUCE
; APPLICANT: RUNDLETT, STEPHEN
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR NON-TARGETED ACTIVATION OF
; TITLE OF INVENTION: ENDOGENOUS GENES
; FILE REFERENCE: 0221-0003C
; CURRENT APPLICATION NUMBER: US/09/479,122
; CURRENT FILING DATE: 2000-01-07
; PRIOR APPLICATION NUMBER: 09/276,820
; PRIOR FILING DATE: 1999-03-26
; PRIOR APPLICATION NUMBER: 09/159,643
; PRIOR FILING DATE: 1998-09-24
; PRIOR APPLICATION NUMBER: 08/941,223
; PRIOR FILING DATE: 1997-09-26
; PRIOR APPLICATION NUMBER: 09/263,814
; PRIOR FILING DATE: 1999-03-08
; PRIOR APPLICATION NUMBER: 09/253,022
; PRIOR FILING DATE: 1999-02-19
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 21
; LENGTH: 5382
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (890)
; OTHER INFORMATION: a, c, t, g, other or unknown
; NAME/KEY: modified_base
; LOCATION: (1042)
; OTHER INFORMATION: a, c, t, g, other or unknown
US-09-479-122-21

Query Match 78.9%; Score 15; DB 4; Length 5382;
Best Local Similarity 100.0%; Pred. No. 1.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 899 GTGCAGGACGTGACA 913

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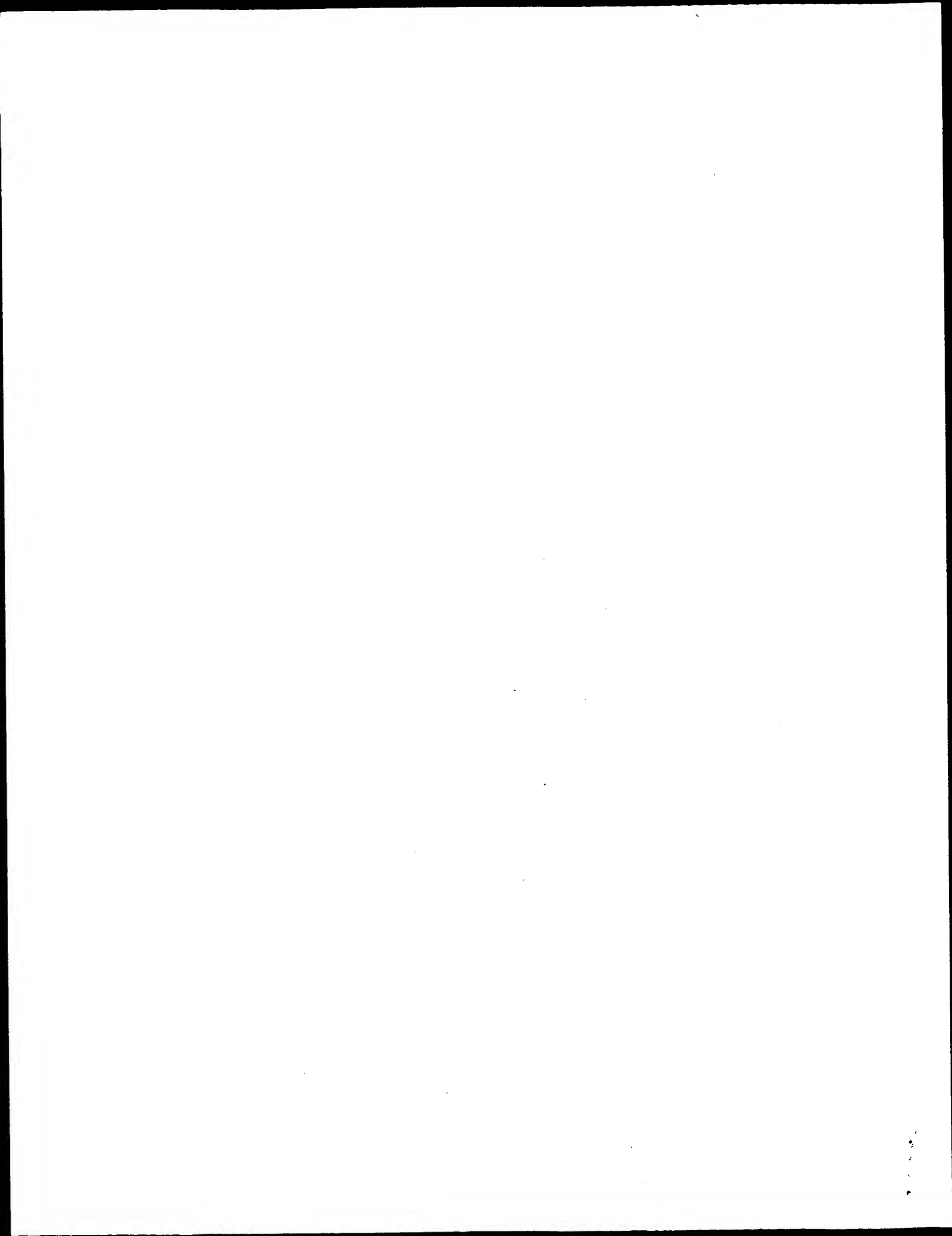
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; Sequence 34, Application US/08646538
; Patent No. 6027881
; GENERAL INFORMATION:
; APPLICANT: Pavlakis, George N.
; APPLICANT: Gaitanaris, George A.
; APPLICANT: Stauber, Roland H.
; APPLICANT: Vournakis, John N.
; TITLE OF INVENTION: Mutant Aequorea victoria Fluorescent
; PROTEIN HAVING INCREASED CELLULAR FLUORESCENCE
; NUMBER OF SEQUENCES: 37
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, 8th Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC Compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/646,538
; FILING DATE: No. 6027881 yet assigned
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weber, Kenneth A.
; REGISTRATION NUMBER: 31,677
; REFERENCE/DOCKET NUMBER: 015280-249000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 34:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7617 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; FEATURE:
; NAME/KEY:
; LOCATION: 1..7617
; OTHER INFORMATION: /note= "pGen-PKGfo25RO"
US-08-646-538-34

Query Match 78.98; Score 15; DB 3; Length 7617;
Best Local Similarity 100.0%; Pred. No. 1.2e-02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 5 GTGCAGGACGTGACA 19
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Db 4186 GTGCAGGACGTGACA 4172

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Job time : 976.413 secs



GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

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(without alignments)  
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Title: US-09-787-562-2

Perfect score: 19

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Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	15	78.9	943	10	US-09-833-381-1259
3	15	78.9	4768	9	US-10-087-523-1
4	15	78.9	4768	10	US-09-816-790-1
5	15	78.9	4768	10	US-09-861-077-1
6	15	78.9	4768	10	US-09-815-825-1
7	15	78.9	4768	10	US-09-815-935-1
8	15	78.9	4768	10	US-09-815-944-1
9	15	78.9	6355	9	US-10-087-523-2
10	15	78.9	6355	10	US-09-816-790-2
11	15	78.9	6355	10	US-09-815-825-2
12	15	78.9	6355	10	US-09-815-935-2
13	15	78.9	6355	10	US-09-815-944-2
14	15	78.9	8388	10	US-09-987-601-1
15	15	78.9	296	10	US-09-294-0938-5837
16	14.8	77.9	729	9	US-09-738-626-1680
17	14.8	77.9	1797	9	US-09-738-626-1682
18	14.8	77.9	1797	9	US-09-738-626-1682
19	14.8	77.9	2748	10	US-09-822-849A-234

c 20	14.8	77.9	6854	9	US-09-922-683-7	Sequence 7, Appl
21	14.8	77.9	43950	12	US-10-060-332-3	Sequence 3, Appl
c 22	14.8	77.9	152331	9	US-10-095-407-16	Sequence 16, Appl
23	14.8	77.9	176373	9	US-10-095-407-17	Sequence 17, Appl
c 24	14.4	75.8	260	10	US-09-864-761-17434	Sequence 17434, A
25	14.4	75.8	338	10	US-09-770-791-871	Sequence 871, App
26	14.4	75.8	392	10	US-09-960-352-3310	Sequence 3310, Ap
27	14.4	75.8	432	10	US-09-960-352-6892	Sequence 6892, Ap
c 28	14.4	75.8	452	10	US-09-864-761-683	Sequence 643, App
29	14.4	75.8	1249	10	US-09-770-445-30	Sequence 30, Appl
30	14.4	75.8	1755	9	US-09-938-842A-1414	Sequence 1414, Ap
c 31	14.2	74.7	226	12	US-10-002-600-63	Sequence 63, Appl
32	14.2	74.7	270	10	US-09-878-574-12415	Sequence 12415, A
33	14.2	74.7	279	10	US-09-878-574-13168	Sequence 13168, A
c 34	14.2	74.7	425	10	US-09-983-965-2423	Sequence 2423, Ap
c 35	14.2	74.7	538	10	US-09-998-598-458	Sequence 458, App
36	14.2	74.7	578	10	US-09-974-300-7790	Sequence 7790, Ap
c 37	14.2	74.7	912	10	US-09-974-300-311	Sequence 311, App
38	14.2	74.7	1098	10	US-09-954-456-2037	Sequence 2037, Ap
39	14.2	74.7	1098	10	US-09-954-456-2070	Sequence 2070, Ap
40	14.2	74.7	1125	12	US-10-044-090-829	Sequence 829, App
41	14.2	74.7	1183	12	US-10-044-090-847	Sequence 847, App
42	14.2	74.7	1353	10	US-09-815-242-9399	Sequence 9399, Ap
43	14.2	74.7	1929	10	US-09-764-898-57	Sequence 57, Appl
44	14.2	74.7	6336	10	US-09-964-824A-114	Sequence 114, App
45	14.2	74.7	6336	10	US-09-880-107-1537	Sequence 1537, Ap

#### ALIGNMENTS

##### RESULT 1

US-09-974-300-8244  
; Sequence 8244, Application US/09974300  
; Patent No. US20020146721A1  
; GENERAL INFORMATION:  
; APPLICANT: Berka, Randy M.  
; APPLICANT: Clausen, Ib Groth  
; TITLE OF INVENTION: Methods For Monitoring Multiple Gene  
; TITLE OF INVENTION: Expression  
; FILE REFERENCE: 10085.500-US  
; CURRENT APPLICATION NUMBER: US/09/974,300  
; CURRENT FILING DATE: 2001-10-05  
; PRIOR APPLICATION NUMBER: 09/680,598  
; PRIOR FILING DATE: 2000-10-06  
; PRIOR APPLICATION NUMBER: 60/279,526  
; PRIOR FILING DATE: 2001-03-27  
; NUMBER OF SEQ ID NOS: 8481  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 8244  
; LENGTH: 461  
; TYPE: DNA  
; ORGANISM: Bacillus clausii  
US-09-974-300-8244

Query Match 83.2%; Score 15.8; DB 10; Length 461;  
Best Local Similarity 89.5%; Pred No. 27;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTCGGTGCAGGACGTGACA 19

Db 9 GTCGGTGCAGGACGTGACA 27

##### RESULT 2

US-09-833-381-1259/c  
; Sequence 1259, Application US/09833381  
; Patent No. US20020132090A1  
; GENERAL INFORMATION:  
; APPLICANT: Robison, Keith E.  
; TITLE OF INVENTION: No. US20020132090A1 Nucleic Acid and Protein Homologs  
; FILE REFERENCE: 5800-119  
; CURRENT APPLICATION NUMBER: US/09/833,381

Mon Jan 6 15:20:23 2003

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; CURRENT FILING DATE: 2001-04-11
; PRIOR APPLICATION NUMBER: 09/516,448
; PRIOR FILING DATE: 2000-02-29
; NUMBER OF SEQ ID NOS: 2050
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1259
; LENGTH: 943
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(943)
; OTHER INFORMATION: n = A,T,C or G
US-09-833-381-1259

Query Match          78.9%; Score 15; DB 10; Length 943;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
   |||||
Db 153 GTGCAGGACGTGACA 139

RESULT 3
US-10-087-523-1
; Sequence 1, Application US/10087523
; Publication No. US20020197624A1
; GENERAL INFORMATION:
; APPLICANT: Klein, Robert D.
; APPLICANT: Brennan, Thomas J.
; TITLE OF INVENTION: METHODS OF CREATING CONSTRUCTS USEFUL FOR INTRODUCING
; TITLE OF INVENTION: SEQUENCES INTO EMBRYONIC STEM CELLS
; FILE REFERENCE: 376472000200
; CURRENT APPLICATION NUMBER: US/10/087,523
; CURRENT FILING DATE: 2002-02-28
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/193,834
; PRIOR FILING DATE: EARLIER FILING DATE: 1998-11-17
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1
; LENGTH: 4768
; TYPE: DNA
; ORGANISM: plasmid vector
; OTHER INFORMATION:
US-10-087-523-1

Query Match          78.9%; Score 15; DB 9; Length 4768;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
   |||||
Db 2824 GTGCAGGACGTGACA 2838

RESULT 4
US-09-816-790-1
; Sequence 1, Application US/09816790
; Patent No. US20020022255A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; APPLICANT: Phillips, Russell
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING
; TITLE OF INVENTION: SULFOTRANSFERASE GENE DISRUPTIONS
; FILE REFERENCE: R-855
; CURRENT APPLICATION NUMBER: US/09/816,790
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,240
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/204,230
; PRIOR FILING DATE: 2000-05-15
; PRIOR APPLICATION NUMBER: US 60/223,173
; PRIOR FILING DATE: 2000-08-07

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; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 4768
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-816-790-1

Query Match          78.9%; Score 15; DB 10; Length 4768;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
   |||||
Db 2824 GTGCAGGACGTGACA 2838

RESULT 5
US-09-861-077-1
; Sequence 1, Application US/09861077
; Patent No. US20020023275A1
; GENERAL INFORMATION:
; APPLICANT: Leviten, Michael W.
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MATRIX
; TITLE OF INVENTION: METALLOPROTEASE GENE DISRUPTIONS
; FILE REFERENCE: R-15
; CURRENT APPLICATION NUMBER: US/09/861,077
; CURRENT FILING DATE: 2000-05-17
; PRIOR APPLICATION NUMBER: US 60/204,972
; PRIOR FILING DATE: 2000-05-17
; PRIOR APPLICATION NUMBER: US 60/215,394
; PRIOR FILING DATE: 2000-06-29
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 4768
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-861-077-1

Query Match          78.9%; Score 15; DB 10; Length 4768;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
   |||||
Db 2824 GTGCAGGACGTGACA 2838

RESULT 6
US-09-815-825-1
; Sequence 1, Application US/09815825
; Patent No. US20020026652A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; APPLICANT: Phillips, Russell
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING CGMP
; TITLE OF INVENTION: PHOSPHODIESTERASE GENE DISRUPTIONS
; FILE REFERENCE: R-849
; CURRENT APPLICATION NUMBER: US/09/815,825
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,142
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/204,227
; PRIOR FILING DATE: 2000-05-15
; PRIOR APPLICATION NUMBER: US 60/216,765
; PRIOR FILING DATE: 2000-07-06
; PRIOR APPLICATION NUMBER: US 60/219,182
; PRIOR FILING DATE: 2000-07-19

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; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 4768
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-825-1

Query Match      78.9%; Score 15; DB 10; Length 4768;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 2824 GTGCAGGACGTGACA 2838

RESULT 7
US-09-815-935-1
; Sequence 1, Application US/09815935
; Patent No. US20020038466A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MAGNESIUM
; TITLE OF INVENTION: DEPENDENT PROTEIN PHOSPHATASE GENE DISRUPTIONS
; FILE REFERENCE: R-723
; CURRENT APPLICATION NUMBER: US/09/815,935
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,235
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/216,249
; PRIOR FILING DATE: 2000-07-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 4768
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-935-1

Query Match      78.9%; Score 15; DB 10; Length 4768;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 2824 GTGCAGGACGTGACA 2838

RESULT 8
US-09-815-944-1
; Sequence 1, Application US/09815944
; Patent No. US20020038467A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; APPLICANT: Matthews, William
; APPLICANT: Moore, Mark
; APPLICANT: Phillips, Russell
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MELANOCYTE
; TITLE OF INVENTION: STIMULATING HORMONE RECEPTOR GENE DISRUPTIONS
; FILE REFERENCE: R-654
; CURRENT APPLICATION NUMBER: US/09/815,944
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,236
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/215,214
; PRIOR FILING DATE: 2000-06-29
; PRIOR APPLICATION NUMBER: US 60/218,075
; PRIOR FILING DATE: 2000-07-12

Query Match      78.9%; Score 15; DB 10; Length 4768;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 2824 GTGCAGGACGTGACA 2838

RESULT 9
US-10-087-523-2
; Sequence 2, Application US/10087523
; Publication No. US20020197624A1
; GENERAL INFORMATION:
; APPLICANT: Klein, Robert D.
; APPLICANT: Brennan, Thomas J.
; TITLE OF INVENTION: METHODS OF CREATING CONSTRUCTS USEFUL FOR INTRODUCING
; TITLE OF INVENTION: SEQUENCES INTO EMBRYONIC STEM CELLS
; FILE REFERENCE: 376472000200
; CURRENT APPLICATION NUMBER: US/10/087,523
; CURRENT FILING DATE: 2002-02-28
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/193,834
; PRIOR FILING DATE: EARLIER FILING DATE: 1998-11-17
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 6355
; TYPE: DNA
; ORGANISM: Plasmid vector
US-10-087-523-2

Query Match      78.9%; Score 15; DB 9; Length 6355;
Best Local Similarity 100.0%; Pred. No. 85;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
Db 4411 GTGCAGGACGTGACA 4425

RESULT 10
US-09-816-790-2
; Sequence 2, Application US/09816790
; Patent No. US20020022255A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; APPLICANT: Phillips, Russell
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING
; TITLE OF INVENTION: SULFOTRANSFERASE GENE DISRUPTIONS
; FILE REFERENCE: R-855
; CURRENT APPLICATION NUMBER: US/09/816,790
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,240
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/204,230
; PRIOR FILING DATE: 2000-05-15
; PRIOR APPLICATION NUMBER: US 60/223,173
; PRIOR FILING DATE: 2000-08-07
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
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us-09-787-562-2.rnpb

Mon Jan 6 15:20:23 2003

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; LENGTH: 6355
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-825-2

Query Match          78.9%; Score 15; DB 10; Length 6355;
Best Local Similarity 100.0%; Pred. No. 85;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
    |||||
Db 4411 GTGCAGGACGTGACA 4425

RESULT 13
US-09-815-935-2
; Sequence 2, Application US/09815935
; Patent No. US20020038466A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MAGNESIUM
; TITLE OF INVENTION: DEPENDENT PROTEIN PHOSPHATASE GENE DISRUPTIONS
; FILE REFERENCE: R-723
; CURRENT APPLICATION NUMBER: US/09/815,935
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,235
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/216,249
; PRIOR FILING DATE: 2000-07-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 6355
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Phage vector
US-09-815-935-2

Query Match          78.9%; Score 15; DB 10; Length 6355;
Best Local Similarity 100.0%; Pred. No. 85;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTGCAGGACGTGACA 19
    |||||
Db 4411 GTGCAGGACGTGACA 4425

RESULT 14
US-09-815-944-2
; Sequence 2, Application US/09815944
; Patent No. US20020038467A1
; GENERAL INFORMATION:
; APPLICANT: Allen, Keith D.
; APPLICANT: Matthews, William
; APPLICANT: Moore, Mark
; APPLICANT: Phillips, Russell
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MELANOCYTE
; TITLE OF INVENTION: STIMULATING HORMONE RECEPTOR GENE DISRUPTIONS
; FILE REFERENCE: R-654
; CURRENT APPLICATION NUMBER: US/09/815,944
; CURRENT FILING DATE: 2001-03-22
; PRIOR APPLICATION NUMBER: US 60/191,236
; PRIOR FILING DATE: 2000-03-22
; PRIOR APPLICATION NUMBER: US 60/215,214
; PRIOR FILING DATE: 2000-06-29
; PRIOR APPLICATION NUMBER: US 60/218,075
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/219,167
; PRIOR FILING DATE: 2000-07-19
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
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; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 2  
; LENGTH: 6355  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Phage vector  
US-09-815-944-2

Query Match 78.9%; Score 15; DB 10; Length 6355;  
Best Local Similarity 100.0%; Pred. No. 85;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTCCAGGACGTGACA 19  
|||||  
Db 4411 GTCCAGGACGTGACA 4425

## RESULT 15

US-09-987-601-1  
; Sequence 1, Application US/09987601  
; Patent No. US20020098223A1  
; GENERAL INFORMATION:  
; APPLICANT: MOULLIER, Phillippe  
; APPLICANT: DANOS, Olivier  
; APPLICANT: HEARD, Jean-Michel  
; APPLICANT: FERRY, Nicholas  
; TITLE OF INVENTION: BIOCOMPATIBLE IMPLANT FOR THE EXPRESSION AND IN VIVO  
; FILE REFERENCE: 0660-0145-0DIV  
; CURRENT APPLICATION NUMBER: US/09/987,601  
; CURRENT FILING DATE: 2001-11-15  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/225,509  
; PRIOR FILING DATE: EARLIER FILING DATE: 1999-01-06  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/523,814  
; PRIOR FILING DATE: EARLIER FILING DATE: 1996-01-19  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: FR 93/04700  
; PRIOR FILING DATE: EARLIER FILING DATE: 1993-04-21  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: FR 93/09185  
; PRIOR FILING DATE: EARLIER FILING DATE: 1993-07-26  
; NUMBER OF SEQ ID NOS: 1  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 1  
; LENGTH: 8388  
; TYPE: DNA  
; ORGANISM: mus musculus, Mo-MuLV, and other  
US-09-987-601-1

Query Match 78.9%; Score 15; DB 10; Length 8388;  
Best Local Similarity 100.0%; Pred. No. 86;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GTCCAGGACGTGACA 19  
|||||  
Db 2238 GTCCAGGACGTGACA 2252

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Job time : 8.41325 secs





GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:41 ; Search time 79.9968 seconds  
(without alignments)  
1281.345 Million cell updates/sec

Title: US-09-787-562-9

Perfect score: 237

Sequence: 1 gctagagtcgtcaggacgt.....cgaggcgctcgccctctg 237

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 381593 seqs, 216252194 residues

Total number of hits satisfying chosen parameters: 763186

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

#### Database :

Published Applications NA:\*

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- 3: /cgn2\_6/ptodata/1/pubpna/US06\_NEW\_PUB.seq:\*
- 4: /cgn2\_6/ptodata/1/pubpna/US06\_PUBCOMB.seq:\*
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- 12: /cgn2\_6/ptodata/1/pubpna/US10\_PUBCOMB.seq:\*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	142.4	60.1	309	10	US-09-965-703-64
2	142.4	60.1	633	9	US-09-925-664-6
3	142.4	60.1	847	10	US-09-950-374-2
4	142.4	60.1	850	10	US-09-950-374-1
5	142.4	60.1	4279	10	US-09-956-988A-1
6	142.4	60.1	4665	10	US-09-759-960-7
7	142.4	60.1	5141	10	US-09-924-859A-9
8	142.4	60.1	5332	8	US-08-961-888-40
9	142.4	60.1	5865	12	US-10-098-035-3
10	142.4	60.1	6620	8	US-08-786-531B-3
11	142.4	60.1	6827	10	US-09-982-610-17
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13	142.4	60.1	7127	7	US-08-779-457-46
14	142.4	60.1	7607	10	US-09-982-610-19
15	142.4	60.1	8578	10	US-09-828-825-1
16	142.4	60.1	8578	10	US-09-828-825-3
17	142.4	60.1	8608	10	US-09-828-825-7
18	142.4	60.1	8623	10	US-09-828-825-5
19	142.4	60.1	8629	10	US-09-828-825-15

20	142.4	60.1	8638	10	US-09-828-825-9	Sequence 9, Appli
21	142.4	60.1	8644	10	US-09-828-825-13	Sequence 13, Appli
22	142.4	60.1	8659	10	US-09-828-825-11	Sequence 11, Appli
23	142.4	60.1	8902	10	US-09-729-418A-1	Sequence 1, Appli
24	142.4	60.1	9108	10	US-09-982-610-45	Sequence 45, Appli
25	142.4	60.1	11152	9	US-09-847-101B-12	Sequence 12, Appli
26	142.4	60.1	11600	9	US-09-847-101B-35	Sequence 35, Appli
27	142.4	60.1	14455	9	US-09-847-101B-15	Sequence 15, Appli
28	142.4	60.1	32480	9	US-09-847-101B-23	Sequence 23, Appli
c 29	141.8	59.8	9077	10	US-09-734-300-1	Sequence 1, Appli
c 30	141.8	59.8	9077	10	US-09-734-300-3	Sequence 3, Appli
31	140.8	59.4	259	10	US-09-919-580-315	Sequence 315, App
32	140.8	59.4	259	10	US-09-919-580-621	Sequence 621, App
33	140.8	59.4	4639	10	US-09-804-481-1	Sequence 1, Appli
34	140.8	59.4	5070	10	US-09-795-006A-41	Sequence 41, Appli
35	140.8	59.4	5432	10	US-09-794-975-9	Sequence 9, Appli
36	140.8	59.4	5446	9	US-09-559-874-5	Sequence 5, Appli
37	140.8	59.4	5446	10	US-09-844-645-3	Sequence 3, Appli
38	140.8	59.4	5458	10	US-09-912-436-11	Sequence 11, Appli
39	140.8	59.4	5458	10	US-09-912-436-12	Sequence 12, Appli
40	140.8	59.4	5614	10	US-09-912-436-7	Sequence 7, Appli
41	140.8	59.4	5614	10	US-09-912-436-8	Sequence 8, Appli
42	140.8	59.4	5651	10	US-09-780-933-6	Sequence 6, Appli
43	140.8	59.4	5695	10	US-09-912-436-9	Sequence 9, Appli
44	140.8	59.4	5695	10	US-09-912-436-10	Sequence 10, Appli
45	140.8	59.4	5864	9	US-09-971-980-1	Sequence 1, Appli

#### ALIGNMENTS

#### RESULT 1

US-09-965-703-64  
; Sequence 64, Application US/09965703  
; Patent No. US20020119521A1  
; GENERAL INFORMATION:  
; APPLICANT: Rohm and Haas Company  
; APPLICANT: Palli, Subba Reddy  
; APPLICANT: Kapitskaya, Marianna Zinovjevna  
; APPLICANT: Cress, Dean Ervin  
; TITLE OF INVENTION: No. US20020119521A1el Ecdysone Receptor-Based Inducible Gene E  
; FILE REFERENCE: A01020B  
; CURRENT APPLICATION NUMBER: US/09/965,703  
; CURRENT FILING DATE: 2001-09-26  
; PRIOR APPLICATION NUMBER: 60/191,355  
; PRIOR FILING DATE: 2000-03-22  
; PRIOR APPLICATION NUMBER: 60/269,799  
; PRIOR FILING DATE: 2001-02-20  
; PRIOR APPLICATION NUMBER: PCT/US01/09050  
; PRIOR FILING DATE: 2001-03-21  
; NUMBER OF SEQ ID NOS: 75  
; SOFTWARE: Patentln version 3.1  
; SEQ ID NO 64  
; LENGTH: 309  
; TYPE: DNA  
; ORGANISM: Simian virus 40  
; FEATURE:  
; NAME/KEY: misc.feature  
; OTHER INFORMATION: No. US20020119521A1el Sequence  
US-09-965-703-64

Query Match	60.1%	Score 142.4;	DB 10;	Length 309;
Best Local Similarity	99.3%;	Pred. No. 6.2e-35;		
Matches 143;	Conservative	0;	Mismatches	1;
			Indels	0;
			Gaps	0;
Qy	94	TGCATCTCAATAGTCAGCACCACATAGTCGCGCCCTTAACCTCCGCCCATCCGCCCTAA	153	
Db	121	TGCATCTCAATAGTCAGCACCACATAGTCGCGCCCTTAACCTCCGCCCATCCGCCCTAA	180	
Qy	154	CTCCGCCAGTTCGCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCTACTAAATTTTATTTATTCGAC	213	
Db	181	CTCCGCCAGTTCGCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCTACTAAATTTTATTTATTCGAC	240	

QY 214 AGCGCGAGGCGCCTCGGCTCTG 237  
 Db 241 AGCGCGAGGCGCCTCGGCTCTG 264

RESULT 2

US-09-925-664-6  
 ; Sequence 6, Application US/09925664  
 ; Patent No. US2002016006A1  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Denney, Jr., Dan W.  
 ; TITLE OF INVENTION: Methods of Treating Lymphoma and Leukemia  
 ; FILE REFERENCE: GENITOPE-06499  
 ; CURRENT APPLICATION NUMBER: US/09/925,664  
 ; CURRENT FILING DATE: 2001-08-09  
 ; PRIOR APPLICATION NUMBER: 09/370,453  
 ; PRIOR FILING DATE: 1999-08-09  
 ; PRIOR APPLICATION NUMBER: 08/644,664  
 ; PRIOR FILING DATE: 1996-05-01  
 ; PRIOR APPLICATION NUMBER: 08/761,277  
 ; PRIOR FILING DATE: 1996-12-06  
 ; NUMBER OF SEQ ID NOS: 80  
 ; SOFTWARE: PatentIn version 3.1  
 ; SEQ ID NO 6  
 ; LENGTH: 633  
 ; TYPE: DNA  
 ; ORGANISM: SR alpha promoter  
 US-09-925-664-6

Query Match 60.1%; Score 142.4; DB 9; Length 633;  
 Best Local Similarity 99.3%; Pred. No. 7.7e-35; Indels 0; Gaps 0;  
 Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCATCCGCCCTAA 153  
 Db 151 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCATCCGCCCTAA 210  
 QY 154 CTCGCCCGAGTTCGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTTATGCGAG 213  
 Db 211 CTCGCCCGAGTTCGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTTATGCGAG 270

RESULT 3

US-09-950-374-2  
 ; Sequence 2, Application US/09950374  
 ; Patent No. US20020141981A1  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Lawman, Patricia  
 ; TITLE OF INVENTION: Materials and Methods for Treating Oncological Disease  
 ; FILE REFERENCE: MOR-200XCD1  
 ; CURRENT APPLICATION NUMBER: US/09/950,374  
 ; CURRENT FILING DATE: 2001-09-10  
 ; PRIOR APPLICATION NUMBER: US 09/394,226  
 ; PRIOR FILING DATE: 1999-09-13  
 ; PRIOR APPLICATION NUMBER: PCT/US99/00787  
 ; PRIOR FILING DATE: 1999-01-14  
 ; PRIOR APPLICATION NUMBER: US 60/071,497  
 ; PRIOR FILING DATE: 1998-01-14  
 ; NUMBER OF SEQ ID NOS: 6  
 ; SEQ ID NO 2  
 ; LENGTH: 847  
 ; TYPE: DNA  
 ; ORGANISM: Streptococcus  
 US-09-950-374-2

Query Match 60.1%; Score 142.4; DB 10; Length 847;  
 Best Local Similarity 99.3%; Pred. No. 8.5e-35; Indels 0; Gaps 0;  
 Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCATCCGCCCTAA 153  
 Db 143 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCATCCGCCCTAA 202  
 QY 154 CTCGCCCGAGTTCGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTTATGCGAG 213  
 Db 203 CTCGCCCGAGTTCGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTTATGCGAG 262  
 QY 214 AGCGCGAGGCGCCTCGGCTCTG 237  
 Db 263 AGCGCGAGGCGCCTCGGCTCTG 286

RESULT 4

US-09-950-374-1  
 ; Sequence 1, Application US/09950374  
 ; Patent No. US20020141981A1  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Lawman, Patricia  
 ; TITLE OF INVENTION: Materials and Methods for Treating Oncological Disease  
 ; FILE REFERENCE: MOR-200XCD1  
 ; CURRENT APPLICATION NUMBER: US/09/950,374  
 ; CURRENT FILING DATE: 2001-09-10  
 ; PRIOR APPLICATION NUMBER: US 09/394,226  
 ; PRIOR FILING DATE: 1999-09-13  
 ; PRIOR APPLICATION NUMBER: PCT/US99/00787  
 ; PRIOR FILING DATE: 1999-01-14  
 ; PRIOR APPLICATION NUMBER: US 60/071,497  
 ; PRIOR FILING DATE: 1998-01-14  
 ; NUMBER OF SEQ ID NOS: 6  
 ; SEQ ID NO 1  
 ; LENGTH: 850  
 ; TYPE: DNA  
 ; ORGANISM: Streptococcus  
 US-09-950-374-1

Query Match 60.1%; Score 142.4; DB 10; Length 850;  
 Best Local Similarity 99.3%; Pred. No. 8.5e-35; Indels 0; Gaps 0;  
 Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCATCCGCCCTAA 153  
 Db 143 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCATCCGCCCTAA 202  
 QY 154 CTCGCCCGAGTTCGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTTATGCGAG 213  
 Db 203 CTCGCCCGAGTTCGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTTATGCGAG 262  
 QY 214 AGCGCGAGGCGCCTCGGCTCTG 237  
 Db 263 AGCGCGAGGCGCCTCGGCTCTG 286

RESULT 5

US-09-956-998A-1  
 ; Sequence 1, Application US/09956998A  
 ; Patent No. US20020082236A1  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Black Jr., Charles A.  
 ; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR ACTIVATING  
 ; TITLE OF INVENTION: GENES OF INTEREST  
 ; FILE REFERENCE: 5722-2(35722/191928)  
 ; CURRENT APPLICATION NUMBER: US/09/956,998A  
 ; CURRENT FILING DATE: 2001-09-20  
 ; PRIOR APPLICATION NUMBER: 09/446,402  
 ; PRIOR FILING DATE: 1999-12-20  
 ; PRIOR APPLICATION NUMBER: 60/050,772  
 ; PRIOR FILING DATE: 1997-06-25  
 ; NUMBER OF SEQ ID NOS: 19  
 ; SOFTWARE: FastSeq for Windows Version 4.0  
 ; SEQ ID NO 1

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; LENGTH: 4279
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Recombinant Molecule containing multiple cloning
; OTHER INFORMATION: site, kozak sequence, LacZ gene.
; NAME/KEY: misc.feature
; LOCATION: (1)...(64)
; OTHER INFORMATION: Multiple cloning site
; NAME/KEY: misc.feature
; LOCATION: (65)...(79)
; OTHER INFORMATION: Consensus sequence for the "Kozak sequence"
; NAME/KEY: prim.transcript
; LOCATION: (80)...(4279)
; OTHER INFORMATION: Beta galactosidase
US-09-956-998A-1

Query Match
Best Local Similarity 60.1%; Score 142.4; DB 10; Length 4279;
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCGCCGCCCTAAGTCCGCCCATCCCGCCCTAA 153
Db 296 TGCATCTCAATTAGTCAGCAACCATAGTCGCCGCCCTAAGTCCGCCCATCCCGCCCTAA 355

QY 154 CTCGCCCCAGTTCGCCCATCTCTCCGCCCATCGCTGACTAAATTTTATTTATGCGAG 213
Db 356 CTCGCCCCAGTTCGCCCATCTCTCCGCCCATCGCTGACTAAATTTTATTTATGCGAG 415

QY 214 AGGCGGAGCGCGCTCGGCTCTG 237
Db 416 AGGCGGAGCGCGCTCGGCTCTG 439

RESULT 6
US-09-759-960-7
; Sequence 7, Application US/09759960
; Patent No. US2001006639A1
; GENERAL INFORMATION:
; APPLICANT: Urban, Robert G.
; APPLICANT: Chicz, Roman M.
; APPLICANT: Collins, Edward J.
; APPLICANT: Hedley, Mary Lynn
; TITLE OF INVENTION: IMMUNOGENIC PEPTIDES FROM THE HPV E7
; TITLE OF INVENTION: PROTEIN
; NUMBER OF SEQUENCES: 33
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson, P.C.
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: MA
; COUNTRY: US
; ZIP: 02110-2804
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows95
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/759,960
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/169,425
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Fraser, Janis K.
; REGISTRATION NUMBER: 34,819
; REFERENCE/DOCKET NUMBER: 06191/004002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-542-5070
; TELEFAX: 617-543-8906
; TELEX: 200154
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; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 4665 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
US-09-759-960-7

Query Match
Best Local Similarity 60.1%; Score 142.4; DB 10; Length 4665;
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCGCCGCCCTAAGTCCGCCCATCCCGCCCTAA 153
Db 288 TGCATCTCAATTAGTCAGCAACCATAGTCGCCGCCCTAAGTCCGCCCATCCCGCCCTAA 347

QY 154 CTCGCCCCAGTTCGCCCATCTCTCCGCCCATCGCTGACTAAATTTTATTTATGCGAG 213
Db 348 CTCGCCCCAGTTCGCCCATCTCTCCGCCCATCGCTGACTAAATTTTATTTATGCGAG 407

QY 214 AGGCGGAGCGCGCTCGGCTCTG 237
Db 408 AGGCGGAGCGCGCTCGGCTCTG 431

RESULT 7
US-09-924-859A-9
; Sequence 9, Application US/09924859A
; Patent No. US20020137113A1
; GENERAL INFORMATION:
; APPLICANT: Godowski, Paul J.
; APPLICANT: Mark, Melanie R.
; APPLICANT: Sadick, Michael D.
; APPLICANT: Shelton, David L.
; APPLICANT: Wong, Wai Lee Tan
; TITLE OF INVENTION: KINASE RECEPTOR ACTIVATION ASSAY
; FILE REFERENCE: P0854C1P2C1
; CURRENT APPLICATION NUMBER: US/09/924,859A
; CURRENT FILING DATE: 2001-08-08
; PRIOR APPLICATION NUMBER: US/09/417,381
; PRIOR FILING DATE: 1999-10-13
; NUMBER OF SEQ ID NOS: 11
; SEQ ID NO 9
; LENGTH: 5141
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Plasmid
US-09-924-859A-9
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Query Match
Best Local Similarity 60.1%; Score 142.4; DB 10; Length 5141;
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCGCCGCCCTAAGTCCGCCCATCCCGCCCTAA 153
Db 188 TGCATCTCAATTAGTCAGCAACCATAGTCGCCGCCCTAAGTCCGCCCATCCCGCCCTAA 247

QY 154 CTCGCCCCAGTTCGCCCATCTCTCCGCCCATCGCTGACTAAATTTTATTTATGCGAG 213
Db 248 CTCGCCCCAGTTCGCCCATCTCTCCGCCCATCGCTGACTAAATTTTATTTATGCGAG 307

QY 214 AGGCGGAGCGCGCTCGGCTCTG 237
Db 308 AGGCGGAGCGCGCTCGGCTCTG 331
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RESULT 8
US-08-961-888-40
; Sequence 40, Application US/08961888
; Patent No. US20010016351A1
; GENERAL INFORMATION:
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; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 5865
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Retroviral vector derived from Moloney Murine
; OTHER INFORMATION: Leukemia Virus
US-10-098-035-3

Query Match          60.1%; Score 142.4; DB 12; Length 5865;
Best Local Similarity 99.3%; Pred. No. 1.5e-34;
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCCATCCGCCCTAA 153
      |||||||
Db 1641 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCCATCCGCCCTAA 1700

QY 154 CTCGCCCGAGTTCGCCCATCTCTCCGCCCATCTCTCGCTGACTACTATTTTATTATGCGAG 213
      |||||||
Db 1701 CTCGCCCGAGTTCGCCCATCTCTCGGCCCATCTCTCGCTGACTACTATTTTATTATGCGAG 1760

QY 214 AGCGCGAGGCGCGCTCGGCTCTG 237
      |||||||
Db 1761 AGCGCGAGGCGCGCTCGGCTCTG 1784

RESULT 10
US-08-786-531B-3
; Sequence 3, Application US/08786531B
; Patent No. US20020015979A1
; GENERAL INFORMATION:
; APPLICANT: Link, Charles J.
; APPLICANT: Levy, John P.
; APPLICANT: Wang, Suming
; APPLICANT: Seregina, Tatiana
; TITLE OF INVENTION: Vehicles for Stable Transfer of Green
; TITLE OF INVENTION: Fluorescent Protein Gene and Methods of Use for Same
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Zarley, McKee, Thomte, Voorhees & Sease
; STREET: 801 Grand Suite 3200
; CITY: Des Moines
; STATE: Iowa
; COUNTRY: United States
; ZIP: 50309
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/786,531B
; FILING DATE: 21-JAN-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/010371
; FILING DATE: 22-JAN-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Nebel, Heidi S.
; REGISTRATION NUMBER: 37,719
; REFERENCE/DOCKET NUMBER: Hgt1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 515-288-3667
; TELEFAX: 515-288-1338
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6620 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; HYPOTHETICAL: NO

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; APPLICANT: Padgett, Kerstien
; APPLICANT: Sorge, Joseph
; TITLE OF INVENTION: NO. US20010016351alel Vector For Gene Expression
; TITLE OF INVENTION: In Prokaryotic And Eukaryotic Systems
; NUMBER OF SEQUENCES: 40
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Howrey & Simon
; STREET: 1299 Pennsylvania Avenue, N.W.
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20004-2402
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/961,888
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Auerbach, Jeffrey I
; REGISTRATION NUMBER: 32,680
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202 383-7451
; TELEFAX: 202 383-6610
; TELEX:
; INFORMATION FOR SEQ ID NO: 40:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 5532 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
US-08-961-888-40

Query Match          60.1%; Score 142.4; DB 8; Length 5532;
Best Local Similarity 99.3%; Pred. No. 1.5e-34;
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 94 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCCATCCGCCCTAA 153
      |||||||
Db 288 TGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTAACTCCGCCCATCCGCCCTAA 347

QY 154 CTCGCCCGAGTTCGCCCATCTCTCCGCCCATCTCTCGCTGACTACTATTTTATTATGCGAG 213
      |||||||
Db 348 CTCGCCCGAGTTCGCCCATCTCTCGGCCCATCTCTCGCTGACTACTATTTTATTATGCGAG 407

QY 214 AGCGCGAGGCGCGCTCGGCTCTG 237
      |||||||
Db 408 AGCGCGAGGCGCGCTCGGCTCTG 431

RESULT 9
US-10-098-035-3
; Sequence 3, Application US/10098035
; Patent No. US20020141983A1
; GENERAL INFORMATION:
; APPLICANT: University of Southern California
; APPLICANT: Weier, Leslie P.
; APPLICANT: McMillan, Minnie
; TITLE OF INVENTION: CONSTRUCTION AND USE OF GENES ENCODING
; TITLE OF INVENTION: PATHOGENIC EPITOPES FOR TREATMENT OF AUTOIMMUNE DISEASE
; FILE REFERENCE: 13761-703-00 US
; CURRENT APPLICATION NUMBER: US/10/098,035
; CURRENT FILING DATE: 2002-03-14
; PRIOR APPLICATION NUMBER: US/08/654,737
; PRIOR FILING DATE: 1996-05-29
; NUMBER OF SEQ ID NOS: 12

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Query Match 60.1%; Score 142.4; DB 10; Length 6827;

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RESULT 13
US - 08-779-457-46
; Sequence 46, Application US/08779457
; Publication No. US2002019357A1
; GENERAL INFORMATION:
; APPLICANT: Carter, Paul J.
; APPLICANT: Chiang, Nancy Y.
; APPLICANT: Kyung, Jin Kim
; APPLICANT: Matthews, William
; APPLICANT: Rodrigues, Maria L.
; TITLE OF INVENTION: WSX RECEPTOR

```

us-09-787-562-9.rnpb

Mon Jan 6 15:20:25 2003

NUMBER OF SEQUENCES: 51  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genentech, Inc.  
STREET: 460 Point San Bruno Blvd  
CITY: South San Francisco  
STATE: California  
COUNTRY: USA  
ZIP: 94080  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: WinPatIn (Genentech)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/779,457  
FILING DATE: 06/20/96  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/667197  
FILING DATE: 06/20/96  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/585005  
FILING DATE: 01/08/96  
ATTORNEY/AGENT INFORMATION:  
NAME: Lee, Wendy M.  
REGISTRATION NUMBER: 40,378  
REFERENCE/DOCKET NUMBER: P0886P2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415/225-1994  
TELEFAX: 415/952-9881  
TELEX: 910/371-7168  
INFORMATION FOR SEQ ID NO: 46:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 7127 base pairs  
TYPE: Nucleic Acid  
STRANDEDNESS: Double  
TOPOLOGY: Linear  
US-08-779-457-46

Query Match 60.1%; Score 142.4; DB 7; Length 7127;  
Best Local Similarity 99.3%; Pred. No. 1.6e-34;  
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 94 TGCATCTCAATTAGTCAGACACCATAGTCGCCGCCCTTAACCTCCGCCCATCCGCCCTAA 153  
DB 2551 TGCATCTCAATTAGTCAGACACCATAGTCGCCGCCCTTAACCTCCGCCCATCCGCCCTAA 2610  
QY 154 CTCGCCCCAGTTCGCCCCATTCCTCCGCCCATCGCTGACTAAATTTTATTATGCGAG 213  
DB 2611 CTCGCCCCAGTTCGCCCCATTCCTCCGCCCATCGCTGACTAAATTTTATTATGCGAG 2670  
QY 214 AGCGCGAGCGCCCTCGGCCCTG 237  
DB 2671 AGCGCGAGCGCCCTCGGCCCTG 2694

RESULT 14  
US-09-982-610-19  
Sequence 19, Application US/09982610  
Patent No. US20020146420A1  
GENERAL INFORMATION:  
APPLICANT: Genentech, Inc.  
Bennett, Brian D.  
Goeddel, David  
Lee, James M.  
Matthews, William  
Tsai, Siao Ping  
Wood, William I.  
TITLE OF INVENTION: PROTEIN TYROSINE KINASE AGONIST ANTIBODIES  
NUMBER OF SEQUENCES: 45  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genentech, Inc.  
STREET: 460 Point San Bruno Blvd

CITY: South San Francisco  
STATE: California  
COUNTRY: USA  
ZIP: 94080  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: WinPatIn (Genentech)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/982,610  
FILING DATE: 17-Oct-2001  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/446,648  
FILING DATE: 1996-MAY-23  
APPLICATION NUMBER: 08/222616  
FILING DATE: 04-APR-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Lee, Wendy M.  
REGISTRATION NUMBER: 40,378  
REFERENCE/DOCKET NUMBER: P0821P3PCT  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415/225-1994  
TELEFAX: 415/952-9881  
TELEX: 910/371-7168  
INFORMATION FOR SEQ ID NO: 19:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 7607 base pairs  
TYPE: Nucleic Acid  
STRANDEDNESS: Single  
TOPOLOGY: Linear  
SEQUENCE DESCRIPTION: SEQ ID NO: 19:  
US-09-982-610-19

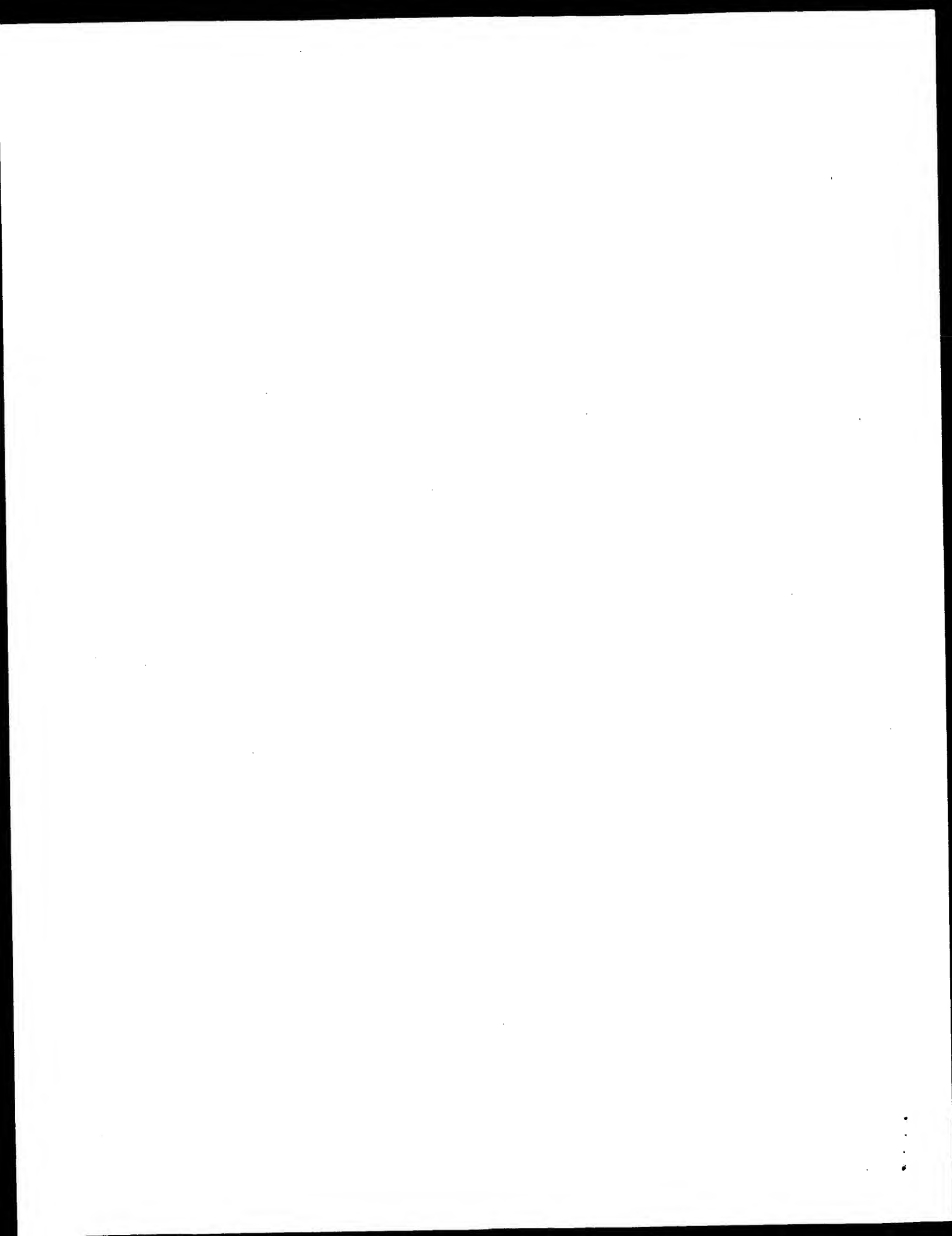
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Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
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DB 4284 TGCATCTCAATTAGTCAGACACCATAGTCGCCGCCCTTAACCTCCGCCCATCCGCCCTAA 4343  
QY 154 CTCGCCCCAGTTCGCCCCATTCCTCCGCCCATCGCTGACTAAATTTTATTATGCGAG 213  
DB 4344 CTCGCCCCAGTTCGCCCCATTCCTCCGCCCATCGCTGACTAAATTTTATTATGCGAG 4403  
QY 214 AGCGCGAGCGCCCTCGGCCCTG 237  
DB 4404 AGCGCGAGCGCCCTCGGCCCTG 4427

RESULT 15  
US-09-828-825-1  
Sequence 1, Application US/09828825  
Patent No. US20020018767A1  
GENERAL INFORMATION:  
APPLICANT: Lee, Seewoo  
APPLICANT: Kim, Han-soo  
TITLE OF INVENTION: Anti-cancer Cellular Vaccine  
FILE REFERENCE: 84906-102  
CURRENT APPLICATION NUMBER: US/09/828,825  
CURRENT FILING DATE: 2001-04-10  
PRIOR APPLICATION NUMBER: KR00-43498  
PRIOR FILING DATE: 2000-07-27  
NUMBER OF SEQ ID NOS: 16  
SOFTWARE: PatentIn Ver. 2.1  
SEQ ID NO 1  
LENGTH: 8578  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
NAME/KEY: enhancer

LOCATION: (1)..(659)  
OTHER INFORMATION: CMV IE  
NAME/KEY: promoter  
LOCATION: (669)..(750)  
OTHER INFORMATION: CMV IE  
NAME/KEY: promoter  
LOCATION: (1067)..(1085)  
OTHER INFORMATION: T7 RNA Promoter  
NAME/KEY: mRNA  
LOCATION: (1090)..(1984)  
OTHER INFORMATION: Human B7.1 (1090-1956 is coding sequence)  
NAME/KEY: RBS  
LOCATION: (2013)..(2593)  
OTHER INFORMATION: IRES sequence  
NAME/KEY: mRNA  
LOCATION: (2627)..(4263)  
OTHER INFORMATION: Human IL12.0 (2640-4223 coding sequence, flexible  
OTHER INFORMATION: linker at 3624-3629)  
NAME/KEY: promoter  
LOCATION: (4352)..(4431)  
OTHER INFORMATION: T3 RNA polymerase promoter  
NAME/KEY: polyA\_signal  
LOCATION: (4362)..(4583)  
OTHER INFORMATION: SV40 fragment containing polyadenylation signal  
NAME/KEY: rep\_origin  
LOCATION: (4678)..(5133)  
OTHER INFORMATION: fl origin of replication  
NAME/KEY: misc\_feature  
LOCATION: (5197)..(6564)  
OTHER INFORMATION: Neo r expression cassette  
NAME/KEY: misc\_feature  
LOCATION: (6975)..(7835)  
OTHER INFORMATION: Ampicillin resistance  
OTHER INFORMATION: Description of Artificial Sequence: Plasmid  
NAME/KEY: CDS  
LOCATION: (2640)..(4223)  
OTHER INFORMATION: IL 12.0 coding sequence - 2 amino acid linker at  
OTHER INFORMATION: 3624  
US-09-828-825-1

Query Match 60.1%; Score 142.4; DB 10; Length 8578;  
Best Local Similarity 99.3%; Pred. No. 1.7e-34;  
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
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QY 154 CTCGCCCGAGTTCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCAT 213  
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Db 5382 CTCGCCCGAGTTCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCAT 5441  
QY 214 AGCGCGAGCGCGCTCGGCTCTG 237  
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Db 5442 AGCGCGAGCGCGCTCGGCTCTG 5465

Search completed: January 4, 2003, 01:06:16  
Job time : 88.9968 secs





GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:46 ; Search time 108.076 Seconds  
(without alignments)  
3231.380 Million cell updates/sec

Title: US-09-787-562-10  
Perfect score: 12  
Sequence: 1 gtcgtcagcga 12

Scoring table: IDENTITY\_NUC  
Gapop 10.0, Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

GenEmbl.\*

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2: gb\_htg.\*  
3: gb\_in.\*  
4: gb\_om.\*  
5: gb\_ov.\*  
6: gb\_pat.\*  
7: gb\_ph.\*  
8: gb\_pi.\*  
9: gb\_pr.\*  
10: gb\_ro.\*  
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12: gb\_sy.\*  
13: gb\_un.\*  
14: gb\_vi.\*  
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16: em\_fun.\*  
17: em\_hum.\*  
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19: em\_mu.\*  
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22: em\_ov.\*  
23: em\_pat.\*  
24: em\_ph.\*  
25: em\_pl.\*  
26: em\_ro.\*  
27: em\_sts.\*  
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33: em\_htg\_mus.\*  
34: em\_htg\_pln.\*  
35: em\_htg\_rod.\*  
36: em\_htg\_mam.\*  
37: em\_htg\_vrt.\*  
38: em\_sy.\*  
39: em\_htgo\_hum.\*  
40: em\_htgo\_mus.\*  
41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	% Match	Query Length	DB ID	Description
1	12	100.0	12	6	AX023668 Sequence
2	12	100.0	24	6	AX023669 Sequence
3	12	100.0	35	6	I58420 Sequence 4
4	12	100.0	36	6	I58427 Sequence 11
5	12	100.0	170	6	E01480 Sequence
6	12	100.0	170	6	E01981 DNA encodin
7	12	100.0	174	6	I58418 Sequence 1
8	12	100.0	177	6	I09058 Sequence 36
9	12	100.0	183	6	A00372 Artificial
10	12	100.0	183	6	A00373 Artificial
11	12	100.0	185	6	E01479 DNA sequenc
12	12	100.0	185	6	E01980 DNA encodin
13	12	100.0	185	6	E02112 DNA sequenc
14	12	100.0	191	6	E02089 DNA encodin
15	12	100.0	191	6	E02091 DNA encodin
16	12	100.0	237	6	AX023667 Sequence
17	12	100.0	269	11	PAU15405
18	12	100.0	296	10	MM031551
19	12	100.0	338	6	E02090 DNA encodin
20	12	100.0	509	1	AF528038 Allothrom
21	12	100.0	515	6	E02113 DNA sequenc
22	12	100.0	575	6	AX401316 Sequence
23	12	100.0	617	9	HS4326857 Homo sapi
24	12	100.0	787	8	AF108439 Coccidiol
25	12	100.0	825	8	AY086953 Arabidops
26	12	100.0	987	3	DME238007 Drosophil
27	12	100.0	1253	1	AF005670 Shevanell
28	12	100.0	1256	1	AF005673 Shevanell
29	12	100.0	1256	1	AF005674 Shevanell
30	12	100.0	1625	1	MTV001 Mycobacte
31	12	100.0	1818	9	BC005158 Homo sapi
32	12	100.0	1845	6	AX123571 Sequence
33	12	100.0	1968	6	AX066951 Sequence
34	12	100.0	1968	6	AX066953 Sequence
35	12	100.0	1985	9	HS805560 Homo sapi
36	12	100.0	2013	9	AK090848 Homo sapi
37	12	100.0	2064	9	AK095929 Homo sapi
38	12	100.0	2087	6	ARI46835 Sequence
39	12	100.0	2103	9	AF189270 Homo sapi
40	12	100.0	2219	1	XORPFCGEN X.oryzae rp
41	12	100.0	2360	8	GPU93077 Gonyalax p
42	12	100.0	2564	9	PATLR3 Papio ham
43	12	100.0	2592	8	AF168786 Sorghum b
44	12	100.0	2624	10	BC020313 Mus muscu
45	12	100.0	2713	1	REU278372 Ralstonia

# ALIGNMENTS

RESULT 1  
AX023668  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL

AX023668  
Sequence 10 from Patent WO0017371.  
AX023668  
AX023668.1 GI:10184029

synthetic construct.  
synthetic construct  
artificial sequences.  
1 (bases 1 to 12)

Binley,K.M. and Naylor,S.  
Polynucleotide constructs and uses thereof  
Patent: WO 0017371-A 10 30-MAR-2000;  
BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD

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Query Match          100.0%; Score 12; DB 6; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
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Db 1 GTCGTGCAGGCA 13

RESULT 4
I58427
LOCUS               36 bp      DNA
DEFINITION          Sequence 11 from patent US 5652120.
ACCESSION            I58427
VERSION              I58427.1 GI:2477665
KEYWORDS
SOURCE               Unknown.
ORGANISM             Unclassified.
REFERENCE            1 (bases 1 to 36)
AUTHORS              Park,S.Kook., Lee,K.Moon., Nho,K.Seung., Koh,Y.Wook., Kwon,C.Hyuk.,
                     Chung,J.Young., Jee,Y.Su. and Yu,Y.Hyo.
TITLE                Gene coding human epidermal growth factor and process for preparing
                     the same
JOURNAL              Patent: US 5652120-A 11 29-JUL-1997;
FEATURES             Location/Qualifiers
source               1..36
/organism="unknown"
BASE COUNT          8 a      9 c      11 g      8 t
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Query Match          100.0%; Score 12; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
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Db 21 GTCGTGCAGGCA 32

RESULT 5
E01480/c
LOCUS               170 bp      DNA
DEFINITION          DNA sequence encoding EGF derivative in recombinant cloning vector.
ACCESSION            E01480
VERSION              E01480.1 GI:2169736
KEYWORDS              JP 1988003791-A/1.
SOURCE               Synthetic construct.
ORGANISM             artificial sequences.
REFERENCE            1 (bases 1 to 170)
AUTHORS              Matsui,A., Harada,Y. and Nakano,T.
TITLE                RECOMBINANT PLASMID
JOURNAL              Patent: JP 1988003791-A 1 08-JAN-1988;
                     HITACHI LTD, HITACHI CHEM CO LTD
COMMENT              OS Artificial gene
                     OC Artificial sequence; Genes.
                     PN JP 1988003791-A/1
                     PD 08-JAN-1988
                     PF 25-JUN-1986 JP 1986146964
                     PI MATSUI AKIKO HARADA YOSHINORI, NAKANO TAKAMORI PC
                     C12N15/00/C07H21/04,C12P21/02,C12P21/02,C12R1.19; CC
                     CC topology: Linear;
                     CC hypothetical: No;
                     CC anti-sense: No;
                     FH Key
                     FH Location/Qualifiers
                     FH CDS
                     FH FT
                     FT RBS
                     FT misc_signal
                     FT 11..16
                     FT /note='XbaI recognition site'.

Query Match          100.0%; Score 12; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
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Db 7 GTCGTGCAGGCA 18

RESULT 3
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LOCUS               35 bp      DNA
DEFINITION          Sequence 4 from patent US 5652120.
ACCESSION            I58420
VERSION              I58420.1 GI:2477658
KEYWORDS
SOURCE               Unknown.
ORGANISM             Unclassified.
REFERENCE            1 (bases 1 to 35)
AUTHORS              Park,S.Kook., Lee,K.Moon., Nho,K.Seung., Koh,Y.Wook., Kwon,C.Hyuk.,
                     Chung,J.Young., Jee,Y.Su. and Yu,Y.Hyo.
TITLE                Gene coding human epidermal growth factor and process for preparing
                     the same
JOURNAL              Patent: US 5652120-A 4 29-JUL-1997;
FEATURES             Location/Qualifiers
source               1..35
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BASE COUNT          7 a      11 c      10 g      7 t
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    Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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  Db 71 GTCGTGCAGGCA 60

  RESULT 6
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  LOCUS      170 bp      RNA      linear      PAT 29-SEP-1997
  DEFINITION DNA encoding epidermal growth factor.
  ACCESSION E01981
  VERSION   E01981.1 GI:2170229
  KEYWORDS JP 1989132383-A/2.
  SOURCE    Unidentified.
  ORGANISM  Homo sapiens
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
    Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  REFERENCE
  AUTHORS    Matsui,A. and Harada,Y.
  TITLE      PLASMID RECOMBINANT
  JOURNAL    Patent: JP 1989132383-A 2 24-MAY-1989;
  COMMENT    HITACHI LTD. HITACHI CHEM CO LTD
  PN  JP 1989132383-A/2
  PD  24-MAY-1989
  PF  18-NOV-1987 JP 1987289332
  PI  MATSUI AKIKO, HARADA YOSHINORI
  PC  C12N15/00,(C12N15/00,C12R1.91);
  CC  strandedness: Double;
  CC  topology: Linear;
  CC  hypothetical: No;
  CC  anti-sense: No;
  CC  *source: clone=pBR322 UG;
  CC  Feature is identified by experimental;
  FH  Key      Location/Qualifiers
  FT  5'UTR      1..17
  FT  CDS      18..167
  FT          /product='epidermal growth factor' FT
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  FT          /product='epidermal growth factor'.
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      /db_xref="taxon:9606"
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    Best Local Similarity 100.0%; Pred. No. 9.4e+03;
    Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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  Db 71 GTCGTGCAGGCA 60

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  LOCUS      174 bp      DNA      linear      PAT 07-OCT-1997
  DEFINITION Sequence 1 from patent US 5652120.
  ACCESSION I58418
  VERSION   I58418.1 GI:2477656
  KEYWORDS

  SOURCE    Unknown.
  ORGANISM  Unclassified.
  REFERENCE 1 (bases 1 to 174)
  AUTHORS    Park,S.Kook., Lee,K.Moon., Nho,K.Seung., Koh,Y.Wook., Kwon,C.Hyuk.,
    Chung,J.Young., Jee,Y.Su. and Yu,Y.Hyo.
  TITLE      Gene coding human epidermal growth factor and process for preparing
    the same
  JOURNAL    Patent: US 5652120-A 1 29-JUL-1997;
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    Best Local Similarity 100.0%; Pred. No. 9.4e+03;
    Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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  Db 54 GTCGTGCAGGCA 43

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  I09058/c
  LOCUS      177 bp      DNA      linear      PAT 02-DEC-1994
  DEFINITION Sequence 36 from Patent WO 8809344.
  ACCESSION I09058
  VERSION   I09058.1 GI:588241
  KEYWORDS
  SOURCE    Unknown.
  ORGANISM  Unclassified.
  REFERENCE 1 (bases 1 to 177)
  AUTHORS    Huston,J.S. and Oppermann,H.
  TITLE      TARGETED MULTIFUNCTIONAL PROTEINS
  JOURNAL    Patent: WO 8809344-A 36 01-DEC-1988;
  FEATURES
    Location/Qualifiers
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    Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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  Db 60 GTCGTGCAGGCA 49

  RESULT 9
  A00372/c
  LOCUS      183 bp      DNA      linear      PAT 04-MAR-1993
  DEFINITION Artificial gene for human epidermal growth factor.
  ACCESSION A00372
  VERSION   A00372.1 GI:344185
  KEYWORDS epidermal growth factor.
  SOURCE    synthetic construct.
  ORGANISM  artificial sequences.
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    /protein_id="CAA00059.1"
    /db_xref="GI:344186"

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PF	25-JUN-1986	JP	1500140302	
PI	HARADA YOSHINORI,	MATSUO AKIKO,	ITO MICHIO	
PC	C12N15/00//C07H21/04.C12P21/02.C12P1/02.C12R1/19			: CC
	strandedness: Double;			
CC	topology: Linear;			
CC	hypothetical: No;			
CC	anti-sense: No;			
FH	key	Location/Qualifiers		
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FT	6..9			
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KEYWORDS      JP 1989257481-A/1.
SOURCE        Homo sapiens.
ORGANISM      Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE     1. (bases 1 to 185)
AUTHORS      Harada,Y., and Matsui,A.
TITLE        DNA HAVING SYNTHETIC GENE FOR PRODUCING HUMAN EPIDERMAL GROWTH
              FACTOR AND PLASMID RECOMBINANT THEREOF
JOURNAL      Patent: JP 1989257481-A 1 13-OCT-1989;
              HITACHI LTD, HITACHI CHEM CO LTD
COMMENT      OS Homo sapiens (Human)
              PN JP 1989257481-A/1
              PD 13-OCT-1989
              PF 08-APR-1988 JP 1988085072
              PI HARADA YOSHINORI, MATSUI AKIKO
              PC C12N15/00;
              CC strandedness: Double;
              CC topology: Linear;
              CC hypothetical: No;
              CC anti-sense: No;
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              FT CDS 18..185
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Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
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Db 71 GTCGTGCAGGCA 60

RESULT 14
E02089/c
LOCUS          191 bp DNA linear PAT 29-SEP-1997
DEFINITION    DNA encoding human epidermal growth factor (hEGF).
ACCESSION     E02089
VERSION       E02089.1 GI:2170331
KEYWORDS      JP 1989247098-A/1.
SOURCE        Synthetic construct.
ORGANISM      artificial sequences.
              1 (bases 1 to 191)
              Harada,Y., Shimizu,N., Fukuzono,S. and Fujimori,K.
              BIOTECHNOLOGICAL PRODUCTION OF HUMAN EPITHELIOCYTE GROWTH FACTOR
              Patent: JP 1989247098-A 1 02-OCT-1989;
              HITACHI LTD, HITACHI CHEM CO LTD
              OS Artificial gene
              OC Artificial sequence: Genes.
              OS Human
              PN JP 1989247098-A/1
              PD 02-OCT-1989
              PF 30-MAR-1988 JP 1988074382
              PI HARADA YOSHINORI, SHIMIZU NORIO, FUKUZONO SHINICHI, PI
              FUJIMORI KIYOSHI
              PC C12P21/02,C07H21/04,C12N15/00,(C12P21/02,C12R1:19); CC
              strandedness: Double;
              CC topology: Linear;
              CC hypothetical: No;
              CC anti-sense: No;
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              CC *source: clone=pbREGF;
              CC Feature is identified by similarity;
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Best Local Similarity 100.0%; Pred. No. 9.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
|||||
Db 71 GTCGTGCAGGCA 60

RESULT 14
E02089/c
LOCUS          191 bp DNA linear PAT 29-SEP-1997
DEFINITION    DNA encoding human epidermal growth factor (hEGF).
ACCESSION     E02089
VERSION       E02089.1 GI:2170331
KEYWORDS      JP 1989247099-A/1.
SOURCE        Synthetic construct.
ORGANISM      artificial sequences.
              1 (bases 1 to 191)
              Shimizu,N., Harada,Y., Fukuzono,S. and Fujimori,K.
              BIOTECHNOLOGICAL PRODUCTION OF HUMAN EPITHELIOCYTE GROWTH FACTOR
              Patent: JP 1989247099-A 1 02-OCT-1989;
              HITACHI LTD, HITACHI CHEM CO LTD
              OS Artificial gene
              OC Artificial sequence: Genes.
              OS Human
              PN JP 1989247099-A/1
              PD 02-OCT-1989
              PF 30-MAR-1988 JP 1988074383
              PI SHIMIZU NORIO, HARADA YOSHINORI, FUKUZONO SHINICHI, PI
              FUJIMORI KIYOSHI
              PC C12P21/02,C07H21/04,C12N15/00,(C12P21/02,C12R1:19); CC
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              CC topology: Linear;
              CC hypothetical: No;
              CC anti-sense: No;
              CC *source: tissue_type=duodenum;
              CC *source: clone=pbREGF;
              CC Feature is identified by similarity;
              FH Key Location/Qualifiers
              FT 5'UTR 24..191
              FT CDS 24..191
              FT /product="human epidermal growth factor"
              FT /db_xref="taxon:32630"
              FT 48 a 45 c 53 g 45 t

FEATURES      source
              Location/Qualifiers
              1..191
              /organism="synthetic construct"
              /db_xref="taxon:32630"
              48 a 45 c 53 g 45 t

BASE COUNT   48 a 45 c 53 g 45 t
ORIGIN
Query Match      100.0%; Score 12; DB 6; Length 191;
Best Local Similarity 100.0%; Pred. No. 9.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
|||||
Db 71 GTCGTGCAGGCA 60

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us-09-787-562-10.rge

Page 6

Db 77 GTCTGCGAGCA 66  
|||||

Search completed: January 3, 2003, 23:54:42  
Job time : 111.076 secs

Result No.	Score	% Match	Length	DB	ID	Description
1	12	100.0	12	21	AAAL2002	Murine HIF-1 space
2	12	100.0	24	21	AAAL2003	Murine HIF-1 space
C 3	12	100.0	35	15	AAQ78668	Human epidermal gr
4	12	100.0	36	15	AAQ78675	Human epidermal gr
C 5	12	100.0	159	20	AAAX8029	Human epidermal gr
C 6	12	100.0	170	10	AAAN90231	Alternative sequen
C 7	12	100.0	174	15	AAQ78658	Epidermal growth f
C 8	12	100.0	177	9	AAAN80185	DNA encoding biosy
C 9	12	100.0	185	9	AAAN81966	Synthetic epiderma

## ALIGNMENTS

XX  
DR WPI; 2000-283595/24.

RESULT 1  
AAA12002

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PT Novel polynucleotide constructs comprising at least two repeats of a  
PT hypoxia response element useful for driving expression of nucleic acids  
PT of interest in a cell under hypoxic conditions  
PS Disclosure: Page 10; 155pp; English.  
XX  
XX This invention describes novel polynucleotide comprising at least 2  
CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible  
CC factor (HIF) consensus binding sites within each of the 2 repeats are  
CC separated by a spacer of at least 20 contiguous nucleotides. The products  
CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic  
CC activity and can be used for gene therapy. The polynucleotides are useful  
CC for delivering nucleic acids of interest to mammalian cells. Lentiviral  
CC vectors are responsive to hypoxic agents and to agents that mimic  
CC hypoxia. This regulation can be harnessed in vivo to enhance the  
CC production of the vector and can be used in vivo to regulate gene  
CC expression in response to a physiological signal. The vectors have  
CC utility in disease, where ischaemia, including hypoxia, is a feature,  
CC e.g. cardiovascular disease, peripheral arterial disease, cancer and  
CC arthritis. The novel regulatory construct is capable of driving very high  
CC levels of transcription under conditions of hypoxia whilst providing only  
CC low basal levels of transcription under normal oxygen conditions. The  
CC polynucleotide construct targets cells within a tumor mass that are under  
CC conditions of hypoxia without affecting normal surrounding tissue. This  
CC sequence represents a murine HIF-1 DNA spacer which is used in the method  
CC of the invention.  
XX  
SQ Sequence 12 BP; 2 A; 3 C; 5 G; 2 T; 0 other;  
Query Match 100.0%; Score 12; DB 21; Length 12;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GTCGTCGAGCA 12  
Db 1 GTCGTCGAGCA 12  
RESULT 2  
AAAL2003  
ID AAA12003 standard; DNA; 24 BP.  
XX  
XX AAA12003;  
AC  
XX 14-AUG-2000 (first entry)  
DT  
XX Murine HIF-1 spacer DNA #2.  
DE  
XX HRE: hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;  
KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;  
KW cardiovascular disease; peripheral arterial disease; cancer; murine; ds.  
KW  
XX  
OS Mus sp.  
XX  
XX WO200017371-A1.  
PN  
XX 30-MAR-2000.  
PD  
XX 22-SEP-1999; 99WO-GB03181.  
PF  
XX 23-SEP-1998; 98WO-GB02885.  
PR 28-JAN-1999; 99GB-0001906.  
PR 16-FEB-1999; 99GB-0003538.  
XX  
XX (OXFO-) OXFORD BIOMEDICA UK LTD.  
PA  
XX Binley KM, Naylor S;  
PI  
XX WPI; 2000-283595/24.  
DR  
XX Novel polynucleotide constructs comprising at least two repeats of a  
PT hypoxia response element useful for driving expression of nucleic acids  
PT of interest in a cell under hypoxic conditions

XX Disclosure: Page 10; 155pp; English.  
PS  
XX This invention describes novel polynucleotide comprising at least 2  
CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible  
CC factor (HIF) consensus binding sites within each of the 2 repeats are  
CC separated by a spacer of at least 20 contiguous nucleotides. The products  
CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic  
CC activity and can be used for gene therapy. The polynucleotides are useful  
CC for delivering nucleic acids of interest to mammalian cells. Lentiviral  
CC vectors are responsive to hypoxic agents and to agents that mimic  
CC hypoxia. This regulation can be harnessed in vivo to enhance the  
CC production of the vector and can be used in vivo to regulate gene  
CC expression in response to a physiological signal. The vectors have  
CC utility in disease, where ischaemia, including hypoxia, is a feature,  
CC e.g. cardiovascular disease, peripheral arterial disease, cancer and  
CC arthritis. The novel regulatory construct is capable of driving very high  
CC levels of transcription under conditions of hypoxia whilst providing only  
CC low basal levels of transcription under normal oxygen conditions. The  
CC polynucleotide construct targets cells within a tumor mass that are under  
CC conditions of hypoxia without affecting normal surrounding tissue. This  
CC sequence represents a murine HIF-1 DNA spacer which is used in the method  
CC of the invention.  
XX  
SQ Sequence 24 BP; 4 A; 5 C; 7 G; 8 T; 0 other;  
Query Match 100.0%; Score 12; DB 21; Length 24;  
Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GTCGTCGAGCA 12  
Db 7 GTCGTCGAGCA 18  
RESULT 3  
AAQ78668/c  
ID AAQ78668 standard; DNA; 35 BP.  
XX  
XX AAQ78668;  
AC  
XX 20-JUL-1995 (first entry)  
DT  
XX Human epidermal growth factor DNA construction oligo, C2.  
DE  
XX Human: epidermal growth factor; hEGF; expression; E. coli;  
KW OmpA leader sequence; universal translation termination;  
KW trpA transcription termination; secretion; ss.  
KW  
XX Synthetic.  
OS  
XX WO9425592-A.  
PN  
XX 10-NOV-1994.  
PD  
XX 25-APR-1994; 94WO-KR00036.  
PF  
XX 26-APR-1993; 93KR-0006978.  
PR 26-APR-1993; 93KR-0006979.  
PR 26-APR-1993; 93KR-0006980.  
XX  
XX (DAEW-) DAEWOONG PHARM CO LTD.  
PA  
XX Chung JY, Jee YS, Koh YW, Kwon CH, Lee KM, Nho KS;  
PI Park SK, Yu YH;  
PI  
XX WPI; 1994-358269/44.  
DR  
XX Novel gene encoding human epidermal growth factor - useful for  
PT transformation of E.coli to give high yield expression.  
PS  
XX Example 2; Fig 2a; 40pp; English.  
XX



CC The sequences given in AAQ78667-76 are oligonucleotides which, when  
 CC ligated together form a human epidermal growth factor (hEGF) coding  
 CC sequence which comprises a HpaI restriction site at its 5' terminal, a  
 CC PstI site at the 3' terminal and Bpu102I, NsiI, MluI, Eco47III and  
 CC AflII restriction sites within the coding sequence. The hEGF contains  
 CC codon usage which is biased for expression in E. coli. This sequence  
 CC may be inserted into a OmpA leader-universal translation termination  
 CC trpA transcription termination sequence between the OmpA leader and  
 CC universal translation termination sequence. This full length sequence  
 CC may then be used for the expression and secretion of hEGF in E. coli.

XX Sequence 35 BP; 7 A; 11 C; 10 G; 7 T; 0 other;

Query Match 100.0%; Score 12; DB 15; Length 35;  
 Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
 |||||  
 Db 24 GTCGTGCAGGCA 13

RESULT 4  
 AAQ78675  
 ID AAQ78675 standard; DNA; 36 BP.  
 AC AAQ78675;

DT 20-JUL-1995 (first entry)

DE Human epidermal growth factor DNA construction oligo, M4.

KW Human; epidermal growth factor; hEGF; expression; E. coli;  
 KW OmpA leader sequence; universal translation termination;  
 KW trpA transcription termination; secretion; ss.

OS Synthetic.

PN WO9425592-A.

PD 10-NOV-1994.

PF 25-APR-1994; 94WO-KR00036.

PR 26-APR-1993; 93KR-0006978.

PR 26-APR-1993; 93KF-0006979.

PR 26-APR-1993; 93KR-0006980.

PA (DAEW-) DAEWOONG PHARM CO LTD.

PI Chung JY, Jee YS, Koh YW, Kwon CH, Lee KM, Nho KS;  
 PI Park SK, Yu YH;

DR WPI; 1994-358269/44.

PT Novel gene encoding human epidermal growth factor - useful for  
 PT transformation of E.coli to give high yield expression.

PS Example 2; Fig 2a; 40pp; English.

CC The sequences given in AAQ78667-76 are oligonucleotides which, when  
 CC ligated together form a human epidermal growth factor (hEGF) coding  
 CC sequence which comprises a HpaI restriction site at its 5' terminal, a  
 CC PstI site at the 3' terminal and Bpu102I, NsiI, MluI, Eco47III and  
 CC AflII restriction sites within the coding sequence. The hEGF contains  
 CC codon usage which is biased for expression in E. coli. This sequence  
 CC may be inserted into a OmpA leader-universal translation termination  
 CC trpA transcription termination sequence between the OmpA leader and  
 CC universal translation termination sequence. This full length sequence  
 CC may then be used for the expression and secretion of hEGF in E. coli.

XX Sequence 36 BP; 8 A; 9 C; 11 G; 8 T; 0 other;

Query Match 100.0%; Score 12; DB 15; Length 36;  
 Best Local Similarity 100.0%; Pred. No. 1.9e+03;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
 |||||  
 Db 21 GTCGTGCAGGCA 32

RESULT 5

AAX89029/C  
 ID AAX89029 standard; DNA; 159 BP.

AC AAX89029;

DT 15-SEP-1999 (first entry)

DE Human epidermal growth factor (hEGF) encoding DNA.

KW Fusion protein; epidermal growth factor; hEGF; human; angiogenin;  
 KW anticancer agent; cancer; cytotoxicity; ss.

OS Homo sapiens.

PN WO923112-A1.

PD 14-MAY-1999.

PF 30-OCT-1998; 98WO-KR00343.

PR 01-NOV-1997; 97KR-0057603.

PA (DAEW-) DAEWOONG PHARM CO LTD.

PI Han S, Kim S, Kim Y, Koo T, Kwon O, Lee B, Park M;  
 PI Park S, Yoon J;

DR WPI; 1999-418417/35.

PD P-PSDB; AAY27102.

PT Human epidermal growth factor and human angiogenin fusion proteins

PS Example 1; Page 36; 73pp; English.

CC The invention provides a fusion protein consisting of human epidermal  
 CC growth factor (hEGF) and human angiogenin joined by a linker. The hEGF-  
 CC angiogenin fusion protein is useful as an anticancer agent. The hEGF-  
 CC tracks down cancer cells expressing hEGF receptors at high level  
 CC following internalisation, and the angiogenin exhibits cytotoxicity by  
 CC degrading ribonucleic acids upon internalisation. The hEGF-angiogenin  
 CC fusion consisting of hEGF and angiogenin, both of which normally exist in  
 CC the human body, exhibit no toxicity following overdose administration.  
 CC The fusion protein can be manufactured in large quantity by cloning in  
 CC bacteria. The fusion protein selectively inhibits the growth of cancer  
 CC cells expressing the hEGF receptor. It does not have a detrimental effect  
 CC on the growth of the normal cells. Further it does not exhibit toxicity  
 CC of conventional chemical anti-cancer agents and it does not cause any  
 CC serious problem by forming antibody against the fusion protein. Targeting  
 CC efficiency is improved using small molecular weight proteins, which are  
 CC 6 kD hEGF and 14.4 kD angiogenin. The present sequence represents a DNA  
 CC encoding a hEGF.

XX Sequence 159 BP; 36 A; 42 C; 47 G; 34 T; 0 other;

Query Match 100.0%; Score 12; DB 20; Length 159;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
 |||||  
 Db 51 GTCGTGCAGGCA 40

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RESULT 6
ID AAN90231/C
XX AAN90231 standard; DNA; 170 BP.
XX AC
XX 01-NOV-1989 (first entry)
XX DT
XX DE Alternative sequence of epidermal growth factor structural gene.
XX PR
XX KW Recombinant plasmid pAMPUG; epidermal growth factor; ampicillin
XX KW resistance; Escherichia coli.
XX PA
XX FH Key Location/Qualifiers
XX FT misc_feature 11..17
XX FT /*tag= a
XX FT CDS 18..164
XX FT /*tag= b
XX FT misc_feature 88..91
XX FT /*tag= c
XX PN JP01132383-A.
XX PD
XX PD 24-MAY-1989.
XX PF 18-NOV-1987; 87JP-0289332.
XX PR 18-NOV-1989; 89JP-0289332.
XX PR (HITA ) HITACHI KK.
XX PA (HITB ) HITACHI CHEMICAL KK.
XX DR WPI; 1989-195592/27.
XX DR P-PSDB; AAP90465.
XX XX
XX PT Recombinant plasmid pAMPUG with epidermal growth activity - comprises the
XX PT epidermal growth factor structural gene joined to HincII site of
XX PT ampicillin resistance gene.
XX PS Disclosure; fig 2; 4pp; Japanese.
XX CC The sequence is an alternative sequence of epidermal growth factor
XX CC structural gene (see AAN90230), which is ligated to the HincII site of
XX CC ampicillin resistance gene, and inserted into recombinant plasmid
XX CC pAMPUG. One or more strong promoter is used, eg tac, lac or trp promoter,
XX CC esp. the amp resistance gene promoter and Shine Dalgarno sequence. The
XX CC EGF is expressed in E.coli. Misc. feature a is an XbaI site, and c is a
XX CC TaqI site. See also AAP90465.
XX SQ Sequence 170 BP; 44 A; 41 C; 45 G; 40 T; 0 other;

Query Match 100.0%; Score 12; DB 10; Length 170;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTCGAGGCA 12
Db |||||
1 171
54 GTCGTCGAGGCA 43

RESULT 8
AAN80185/C
ID AAN80185 standard; DNA; 177 BP.
XX AC
XX AC AAN80185;
XX DT 14-NOV-1990 (first entry)
XX XX
XX DE DNA encoding biosynthetic multifunctional protein.
XX XX Biosynthetic multifunctional protein; biosynthetic antibody binding site;
XX KW protein trailer; epidermal growth factor; ss DNA.
XX CC
XX FH Key Location/Qualifiers
XX FT CDS 1..171
XX FT /*tag= a
XX PN W08809344-A.
XX PD 01-DEC-1988.
XX PD 19-MAY-1988; 88WO-US01737.
XX PF
XX PR 21-MAY-1987; 87US-0052800.
XX XX
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PA (CREA-) CREATIVE BIOMOLECULES INC.
XX
XX PI Huston JS, Oppermann H;
XX
XX WPI: 1988-353928/49.
XX P-PSDB; AAP80159.
XX
XX Recombinant multifunctional protein - having antibody binding site and a
XX sequence for biological activity, ion sequestering or binding to a
XX solid support.
XX
XX Disclosure; ; 115pp; English.
XX
XX The DNA encodes a biosynthetic multifunctional protein including a single
XX chain biosynthetic antibody binding site and an epidermal growth factor
XX protein trailer.
XX
XX Sequence 177 BP; 39 A; 45 C; 53 G; 40 T; 0 other;
XX
XX Query Match 100.0%; Score 12; DB 9; Length 177;
XX Best Local Similarity 100.0%; Pred. No. 1.8e+03;
XX Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 GTCGTGCAGGCA 12
XX |||||||
XX Db 60 GTCGTGCAGGCA 49
XX
XX RESULT 9
XX AAN81966/c
XX ID AAN81966 standard; DNA; 185 BP.
XX
XX AC AAN81966;
XX
XX DT 09-OCT-1990 (first entry)
XX
XX DE Synthetic epidermal growth factor gene.
XX
XX KW Epidermal growth factor; ds.
XX
XX OS Synthetic.
XX
XX FH Key Location/Qualifiers
XX CDS 18..185
XX FT /*tag= a
XX FT /product=EGF
XX FT 6..9
XX FT /*tag= b
XX
XX PN JP63003790-A.
XX
XX PD 08-JAN-1988.
XX
XX PF 25-JUN-1986; 86JP-0146963.
XX
XX PR 25-JUN-1986; 86JP-0146963.
XX
XX (HITA ) HITACHI KK.
XX (HITB ) HITACHI CHEMICAL KK.
XX
XX WPI: 1988-046256/07.
XX P-PSDB; AAP81522.
XX
XX Recombinant plasmid with higher prodn. efficiency - is prepd. by
XX inserting a structural gene, encoding a 53-residue polypeptide,
XX downstream of tac promoter.
XX
XX Disclosure; ; Japanese.
XX
XX The sequence was synthesized from 40 oligonucleotide fragments. The
XX 5' end of the sense strand overhangs the complementary strand by 4
XX bases; the 5' end of the complementary strand overhangs the 3' end of
XX the sense strand by -CTAG. The sequence is also given in J633791-A.
XX
XX Sequence 185 BP; 46 A; 44 C; 52 G; 43 T; 0 other;
XX
XX Query Match 100.0%; Score 12; DB 10; Length 185;
XX Best Local Similarity 100.0%; Pred. No. 1.8e+03;
XX Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 GTCGTGCAGGCA 12
XX |||||||
XX Db 71 GTCGTGCAGGCA 60
XX
XX RESULT 10
XX AAN90230/c
XX ID AAN90230 standard; DNA; 185 BP.
XX
XX AC AAN90230;
XX
XX DT 01-NOV-1989 (first entry)
XX
XX DE Epidermal growth factor structural gene.
XX
XX KW Recombinant plasmid pAMPUG; ampicillin resistance;
XX Escherichia coli.
XX
XX FH Key Location/Qualifiers
XX CDS 11..16
XX FT /*tag= a
XX FT 18..179
XX FT /*tag= b
XX FT misc_feature 88..91
XX FT /*tag= c
XX
XX PN JF01132383-A.
XX
XX PD 24-MAY-1989.
XX
XX PF 18-NOV-1987; 87JP-0289332.
XX
XX PR 18-NOV-1989; 89JP-0289332.
XX
XX (HITA ) HITACHI KK.
XX (HITB ) HITACHI CHEMICAL KK.
XX
XX WPI: 1989-195592/27.
XX P-PSDB; AAP90465.
XX
XX Recombinant plasmid pAMPUG with epidermal growth activity - comprises the
XX epidermal growth factor structural gene joined to HincII site of
XX ampicillin resistance gene.
XX
XX Disclosure; fig 2; 4pp; Japanese.
XX
XX The sequence is of epidermal growth factor structural gene, which is
XX ligated to the HincII site of ampicillin resistance gene, and inserted
XX into recombinant plasmid pAMPUG. One or more strong promoter is used,
XX eg tac, lac or trp promoter, esp. the amp resistance gene promoter and
XX Shine Dalgarno sequence. The EGF is expressed in E.coli (see AAP90464
XX for encoded peptide). Misc. feature a is an XbaI site, and c is a TaqI
XX site. See also AAN90231 and AAP90465.
XX
XX Sequence 185 BP; 46 A; 44 C; 52 G; 43 T; 0 other;
XX
XX Query Match 100.0%; Score 12; DB 10; Length 185;
XX Best Local Similarity 100.0%; Pred. No. 1.8e+03;
XX Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 GTCGTGCAGGCA 12
XX |||||||
XX Db 71 GTCGTGCAGGCA 60
XX
XX RESULT 11
```

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AA091199/c
ID AA091199 standard; DNA; 191 BP.
XX AC AA091199;
XX DT 05-FEB-1990 (first entry)
XX DE Chemically synthesised human epidermal growth factor (hEGF) gene.
XX KW Human epidermal growth factor (hEGF); tryptophan regulatory gene;
XX KW PBR322; pTREBT; pTRLBT
XX FT Key Location/Qualifiers
XX FT CDS 24..191
XX FT /*tag= a
XX PN EP335400-A.
XX PD 04-OCT-1989.
XX PF 30-MAR-1989; 89EP-0105639.
XX PR 30-MAR-1988; 88JP-0074383.
XX PA (HITA ) HITACHI KK.
XX PI Shimizu N, Harada Y, Fukuzono S, Fujimori K;
XX DR WPI; 1989-286891/40.
XX DR P-PSDB;P9187.
XX DT DNA contg. human epidermal growth factor gene
XX PT - associated with E. coli tryptophan operon components
XX PS Claim 1; figure 1; 26pp; English.
XX CC Entire sequence of organically synthesised hEGF gene. Has 5' AATT single-
XX CC stranded sticky end on coding strand and 5' CTAG single-stranded sticky
XX CC end on non-coding strand.
XX SQ Sequence 191 BP; 48 A; 45 C; 53 G; 45 T; 0 other;

Query Match 100.0%; Score 12; DB 10; Length 191;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
DB 77 GTCGTGCAGGCA 66

RESULT 12
AA012001
ID AA012001 standard; DNA; 237 BP.
XX AC AA012001;
XX DT 14-AUG-2000 (first entry)
XX DE Murine PGK HRE derived promoter OBHrell DNA.
XX KW HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotrophic;
XX KW cardiant; cytosstatic; antiarthritic; gene therapy; ischaemia; arthritis;
XX KW cardiovascular disease; peripheral arterial disease; cancer;
XX KW phosphoglycerate kinase; PGK; murine; promoter; OBHrell; ds.
XX OS Mus sp.
XX PN WO200017371-A1.
XX PD 30-MAR-2000.
XX PF 22-SEP-1999; 99WO-GB03181.

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XX 23-SEP-1998; 98WO-GB02885.
XX 28-JAN-1999; 99GB-0001906.
XX 16-FEB-1999; 99GB-0003538.
XX (OXFO-) OXFORD BIOMEDICA UK LTD.
XX PI Binley KM, Naylor S;
XX WPI; 2000-283595/24.
XX Novel polynucleotide constructs comprising at least two repeats of a
XX hypoxia response element useful for driving expression of nucleic acids
XX of interest in a cell under hypoxic conditions
XX Example 1; Page 68; 155pp; English.
XX This invention describes novel polynucleotide comprising at least 2
XX repeats of a hypoxia response element (HRE), where the hypoxia-inducible
XX factor (HIF) consensus binding sites within each of the 2 repeats are
XX separated by a spacer of at least 20 contiguous nucleotides. The products
XX of the invention have vasotropic, cardiant, cytosstatic and antiarthritic
XX activity and can be used for gene therapy. The polynucleotides are useful
XX for delivering nucleic acids of interest to mammalian cells. Lentiviral
XX vectors are responsive to hypoxic agents and to agents that mimic
XX hypoxia. This regulation can be harnessed in vitro to enhance the
XX production of the vector and can be used in vivo to regulate gene
XX expression in response to a physiological signal. The vectors have
XX utility in disease, where ischaemia, including hypoxia, is a feature,
XX e.g. cardiovascular disease, peripheral arterial disease, cancer and
XX arthritis. The novel regulatory construct is capable of driving very high
XX levels of transcription under conditions of hypoxia whilst providing only
XX low basal levels of transcription under normal oxygen conditions. The
XX polynucleotide construct targets cells within a tumor mass that are under
XX conditions of hypoxia without affecting normal surrounding tissue. This
XX sequence represents a murine phosphoglycerate kinase (PGK) HRE derived
XX promoter OBHrell which is described in the method of the invention.
XX SQ Sequence 237 BP; 43 A; 82 C; 56 G; 56 T; 0 other;

Query Match 100.0%; Score 12; DB 21; Length 237;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
DB 31 GTCGTGCAGGCA 42

RESULT 13
AA020579/c
ID AA020579 standard; DNA; 330 BP.
XX AC AA020579;
XX DT 09-JUN-1992 (first entry)
XX DE [Phe23]EGF in pTA 1522-Eco.
XX KW Epidermal growth factor; variant; mutant; receptor; anticancer; ds.
XX OS Homo sapiens.
XX FT Key Location/Qualifiers
XX FT CDS 95..319
XX FT /*tag= a
XX FT /label= EGF
XX FT repeat_unit 4..25
XX FT /*tag= b
XX FT /rpt_type= inverted
XX FT repeat_unit 28..41
XX FT /*tag= c
XX FT /rpt_type= inverted

```

FT repeat\_unit 44...111  
 FT /\*tag= d  
 FT /rpt\_type= inverted  
 FT misc\_signal 43..49  
 FT /\*tag= e  
 FT /label= PB  
 FT RBS 83..86  
 FT /\*tag= f  
 FT sig\_peptide 95..157  
 FT /\*tag= g  
 FT /label= phoA\_signal  
 FT /note= "21 amino acids"  
 FT mat\_peptide 158..319  
 FT /\*tag= h  
 FT /label= EGF  
 FT misc\_difference 224..226  
 FT /\*tag= i  
 FT /note= "mutated to encode Phe instead of Ile"  
 FT JP03294293-A.  
 PN 25-DEC-1991.  
 XX 11-APR-1990; 90JP-0093921.  
 XX 11-APR-1990; 90JP-0093921.  
 PR (YUEI-) YUEI SEIVAKU KK.  
 PA WPI: 1992-052025/07.  
 DR P-PSDB; AAR20623.  
 XX

Human epithelial cell growth factor variant - with higher receptor activity than natural EGF, useful in development of anticancer drugs

PS Disclosure; Fig 1; 6pp; Japanese.

CC pTA1522-Eco comprises a phoA promoter-, EGF- and a phoA- signal sequence. The codon for Ile at position 23 is changed to TTT (encoding Phe) by oligonucleotide site-directed mutagenesis to produce the new variant. The variant has a higher affinity for receptors than natural hEGF and may be used in the development of anticancer drugs.

XX Sequence 330 BP; 83 A; 72 C; 78 G; 97 T; 0 other;

Query Match 100.0%; Score 12; DB 13; Length 330;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12  
 |||||  
 Db 208 GTCGTGCAGGCA 197

RESULT 14  
 AAA44708

ID AAA44708 standard; cDNA; 346 BP.

XX

AC AAA44708;

XX 21-AUG-2000 (first entry)

XX Human secreted expressed sequence tag SEQ ID NO:1283.

XX Human; mouse; chicken; rat; secreted expressed sequence tag; SEST;  
 expressed sequence tag; EST; probe; chemotactic; proliferative;  
 immunomodulatory; haematopoietic; chemokinetic; analgesic; haemostatic;  
 thrombolytic; antiinflammatory; cytostatic; antibacterial; antifungal;  
 antiviral; antidiabetic; antiasthmatic; vulnery; antiparkinsonian;  
 antiulcer; osteopathic; neuroprotective; nootropic; antipsoriatic;  
 cerebroprotective; anticonvulsant; antidepressant; gene therapy;

KW vaccine; autoimmune disorder; multiple sclerosis; allergic condition;  
 KW insulin dependent diabetes; asthma; myeloid cell deficiency; ulcer;  
 KW lymphoid cell deficiency; burn; osteoporosis; osteoarthritis;  
 KW central nervous system disorder; Alzheimer's disease; stroke;  
 KW Parkinson's disease; Huntington's disease; coagulation disorder;  
 KW haemophilia; thrombosis; inflammatory disorder; Crohn's disease;  
 KW tumour; infection; depression; psoriasis; ss.

XX Homo sapiens.

XX WO200021991-A1.

XX 20-APR-2000.

XX 15-OCT-1999; 99WO-US24206.

XX 15-OCT-1998; 98US-0104436.

XX (GEMY ) GENETICS INST INC.

XX Jacobs K, McCoy JM, LaVallie ER, Collins-Racie LA, Evans C;

PI Merberg D, Treacy M, Bowman MR;

XX WPI: 2000-317938/27.

XX Isolated polynucleotides, and encoded proteins, comprising secreted expressed sequence tags (SESTs), useful for treating various disorders such as autoimmune, infectious, and central nervous system disorders -  
 Claim 1; Page 535; 803pp; English.

CC AAA43426 to AAA45925 represent specifically claimed secreted expressed sequence tags (SESTs), isolated from human, mouse, chicken and rat tissue sources. The SESTs can have a range of activities depending on the tissues they were isolated from. The activities include:  
 CC chemokinetic; proliferative; immunomodulatory; haematopoietic;  
 CC cytostatic; antibacterial; antifungal; thrombolytic; antiinflammatory;  
 CC antiasthmatic; vulnery; antiulcer; osteopathic; neuroprotective;  
 CC nootropic; antiparkinsonian; antipsoriatic; cerebroprotective;  
 CC anticonvulsant; and antidepressant. The SESTs can be used for gene therapy and in vaccines. The SESTs are useful as probes for the identification and isolation of full-length cDNAs and genomic DNA molecules which correspond to the SESTs. Proteins encoded by the SESTs are useful in assays for determining biological activity and raising antibodies. They may be useful for treatment of autoimmune disorders (multiple sclerosis, insulin dependent diabetes), allergic conditions (asthma), myeloid or lymphoid cell deficiencies, wounds, burns, ulcers, osteoporosis, osteoarthritis, central nervous system disorders (Alzheimer's, Parkinson's, Huntington's disease, stroke), coagulation disorders (haemophilia, thrombosis), inflammatory disorders (Crohn's disease), tumours, bacterial, fungal or viral infections, depression and psoriasis. AAA45926 to AAA45931 represent linker variants which are given in the exemplification of the present invention.

XX Sequence 346 BP; 79 A; 61 C; 131 G; 75 T; 0 other;

Query Match 100.0%; Score 12; DB 21; Length 346;  
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12  
 |||||

Db 87 GTCGTGCAGGCA 98

RESULT 15

AAAN92328/C

ID AAAN92328 standard; DNA; 515 BP.

XX AC AAAN92328;

XX 25-APR-1990 (first entry)

```

XX Synthetic human epidermal growth factor.
DE Human epidermal growth factor; EGF;
KW Human epidermal growth factor; EGF;
XX
XX Key Location/Qualifiers
PH 18..570
CDS /*tag= a
FT repeat_unit 171..350
FT /*tag= b
XX
XX JP01257482-A.
XX
XX 13-OCT-1989.
XX
XX 08-APR-1988; 88JP-0085073.
XX
XX 08-APR-1988; 88JP-0085073.
XX
XX (HITA ) HITACHI KK.
PA (HITB ) HITACHI CHEMICAL KK.
XX
XX WPI; 1989-344714/47.
DR P-PSDB; AAP93397.
XX
XX DNA contg. synthetic gene - used for human epidermal growth factor prodn.
XX
XX Disclosure; fig. 2; 7pp; Japanese.
XX
XX The gene has the following structure: tag b:(AAPI,m)m)n:
CC Tag b is a repeating unit encoding hEGF. The synthetic gene
CC can be inserted into a plasmid vector for efficient prodn. of EGF.
XX
XX Sequence 515 BP; 125 A; 125 C; 148 G; 117 T; 0 other;
SQ
Query Match 100.0%; Score 12; DB 10; Length 515;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTCGTCAGGCA 12
DB 71 GTCGTCAGGCA 60
Search completed: January 3, 2003, 23:20:44
Job time : 21.0631 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:17 ; Search time 159.558 Seconds  
(without alignments)  
1218.024 Million cell updates/sec

Title: US-09-787-562-10  
Perfect score: 12  
Sequence: 1 gtcgtgcaggca 12

Scoring table: IDENTITY\_NUC  
Gapop 10.0, Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:\*

- 1: em\_estba:\*
- 2: em\_esthum:\*
- 3: em\_estnu:\*
- 4: em\_estnu:\*
- 5: em\_estov:\*
- 6: em\_estpl:\*
- 7: em\_estro:\*
- 8: em\_hic:\*
- 9: gb\_est1:\*
- 10: gb\_est2:\*
- 11: gb\_hic:\*
- 12: gb\_est3:\*
- 13: gb\_est4:\*
- 14: gb\_est5:\*
- 15: em\_estfun:\*
- 16: em\_estom:\*
- 17: gb\_gss:\*
- 18: em\_gss\_hum:\*
- 19: em\_gss\_inv:\*
- 20: em\_gss\_pln:\*
- 21: em\_gss\_vit:\*
- 22: em\_gss\_fun:\*
- 23: em\_gss\_mam:\*
- 24: em\_gss\_mus:\*
- 25: em\_gss\_other:\*
- 26: em\_gss\_pro:\*
- 27: em\_gss\_rod:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	12	100.0	137	12	BG050523 FMI_72_D0
2	12	100.0	174	12	BF837005 PM3-HT090
3	12	100.0	177	9	AI030525 UI-R-CO-1
4	12	100.0	180	10	BE150969 RC4-HT07
5	12	100.0	192	13	BI024277 CM3-MT029
6	12	100.0	212	13	BI034110 PM2-NN016

7	12	100.0	221	9	AA815339
8	12	100.0	239	12	BF760820
9	12	100.0	248	17	BH227331
10	12	100.0	253	13	BM145815
11	12	100.0	256	12	BF256224
12	12	100.0	261	12	BF146813
13	12	100.0	286	10	BB179675
14	12	100.0	292	14	BM820141
15	12	100.0	296	9	AT004345
16	12	100.0	297	12	BE766861
17	12	100.0	298	9	AA469407
18	12	100.0	300	9	AJ467211
19	12	100.0	300	9	AJ468211
20	12	100.0	305	10	BB491289
21	12	100.0	307	10	AW401951
22	12	100.0	316	10	BE604099
23	12	100.0	323	17	AZ577296
24	12	100.0	325	10	BE146189
25	12	100.0	325	12	BE863787
26	12	100.0	328	14	248448
27	12	100.0	330	9	AI867483
28	12	100.0	331	9	AV158985
29	12	100.0	332	9	AA741679
30	12	100.0	332	10	BE517981
31	12	100.0	336	14	BQ166106
32	12	100.0	340	9	AL816780
33	12	100.0	344	14	BQ164373
34	12	100.0	346	13	BI974238
35	12	100.0	357	9	AA666066
36	12	100.0	359	12	BF359469
37	12	100.0	361	10	AW87486
38	12	100.0	363	10	BE086558
39	12	100.0	365	9	AI007372
40	12	100.0	368	10	BE634826
41	12	100.0	369	14	BQ279572
42	12	100.0	373	17	AQ099324
43	12	100.0	375	12	BG013635
44	12	100.0	378	12	BG234973
45	12	100.0	384	9	AA146187

#### ALIGNMENTS

RESULT 1  
BG050523  
LOCUS  
DEFINITION  
FMI\_72\_D03\_g1\_A003 Floral-Induced Meristem 1 (FMI) Sorghum  
PROPINQUUM CDNA, mRNA sequence.  
ACCESSION  
BG050523  
VERSION  
BG050523.1  
KEYWORDS  
EST.  
SOURCE  
Sorghum propinquum.  
ORGANISM  
Sorghum propinquum  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC  
clade; Panicoideae; Andropogoneae; Sorghum.  
REFERENCE  
1 (bases 1 to 137)  
Cordonnier-Pratt, M.-M., Gingle, A., Sudman, M., Marsala, C. and Pratt, L.H.  
TITLE  
An EST database from Sorghum: floral-induced meristems  
JOURNAL  
Unpublished (2000)  
COMMENT  
Contact: Cordonnier-Pratt MM  
Laboratory for Genomics and Bioinformatics  
The University of Georgia, Department of Plant Biology  
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA  
Tel: 706 542 1860  
Fax: 706 583 0210  
Email: mmpratt@uga.edu  
Sequences have been trimmed to exclude PolyA, vector and regions  
below Phred quality 16. The threshold for highest quality sequence  
is 20.  
Seq primer: PolyTMix

High quality sequence start: 29  
High quality sequence stop: 70  
POLYA-No. Location/Qualifiers  
1. .137  
/organism="Sorghum propinquum"  
/db\_xref="taxon:132711"  
/clone\_lib="Floral-Induced Meristem 1 (FM1)"  
/notes="Organ: Floral-Induced meristems; Vector: pBluescript II from Lambda Zap II; Site\_1: XhoI; Site\_2: EcoRI; mature plants were placed in a growth chamber for 15 days with 16 hr darkness and 8 hr light (flowering is induced by short-day conditions); 16 days after being returned to the greenhouse under natural long days during late April/early May, meristems were harvested. The library was made from poly-A RNA in the cloning vector lambda ZAP II. Clones to be sequenced were prepared by mass excision." 34 a 26 c 31 g 46 t

BASE COUNT 38 a 51 c 34 g 51 t  
ORIGIN  
Query Match 100.0%; Score 12; DB 12; Length 174;  
Best Local Similarity 100.0%; Pred. No. 3.6e+04;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GTCGTGCAGGCA 12  
|||||  
Db 9 GTCGTGCAGGCA 20

RESULT 3  
AI030525  
LOCUS  
DEFINITION  
UI-R-C0-ji-c-04-0-UI.s1 UI-R-C0 Rattus norvegicus cDNA clone  
UI-R-C0-ji-c-04-0-UI 3', mRNA sequence.  
AI030525  
ACCESSION  
VERSION  
AI030525.1 GI:3248351  
KEYWORDS  
EST.  
SOURCE  
Norway rat.  
Rattus norvegicus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.

REFERENCE  
1 (bases 1 to 177)  
Bonaldo,M.F., Lennon,G. and Soares,M.B.  
Normalization and subtraction: two approaches to facilitate gene  
discovery  
Genome Res. 6 (9), 791-806 (1996)  
97044477  
Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: mscores@blue.weeg.uiowa.edu  
The sequence tag present in the cDNA between the NotI site and the  
oligo-dT track served to identify it as a clone from the normalized  
adult Spleen library. cDNA library Preparation: M. Fatima Bonaldo,  
Ph.D. Clone distribution: clones will be available through Research  
Genetics This clone is also available through the I.M.A.G.E.  
Consortium at LBNL (info@image.lbnl.gov). IMAGE ID=1783023  
Seq primer: M13 Forward  
POLYA-No.

FEATURES  
source  
1. 177  
/organism="Rattus norvegicus"  
/strain="Sprague-Dawley"  
/db\_xref="taxon:10116"  
/clone="UI-R-C0-ji-c-04-0-UI"  
/clone\_lib="UI-R-C0"  
/dev\_stage="adult"  
/lab\_host="DH10B (Life Technologies)"  
/note="Vector: pMT3p-Pac (Pharmacia) with a modified  
polylinker; Site\_1: Not I; Site\_2: Eco RI; The UI-R-C0  
library is a subtracted library derived from the UI-R-A1  
and UI-R-E1 libraries. The UI-R-A1 library consisted of a  
mixture of individually tagged normalized libraries  
constructed from rat placenta, adult lung, brain, liver,  
kidney, heart, spleen, ovary, and muscle. The UI-R-E1  
library consisted of a mixture of individually tagged  
normalized libraries constructed from 8, 12 and 18-day  
embryo. The tag is a string of 3-5 nucleotides present  
between the Not I site and the oligo-dT track which

FEATURES  
source  
1. 174  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_lib="HT0909"  
/dev\_stage="Adult"

BASE COUNT 34 a 26 c 31 g 46 t  
ORIGIN  
Query Match 100.0%; Score 12; DB 12; Length 137;  
Best Local Similarity 100.0%; Pred. No. 3.5e+04;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GTCGTGCAGGCA 12  
|||||  
Db 6 GTCGTGCAGGCA 17

RESULT 2  
BF837005  
LOCUS  
DEFINITION  
BF837005  
ACCESSION  
VERSION  
BF837005.1 GI:12189164  
KEYWORDS  
EST.  
SOURCE  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 174)  
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,  
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.,  
Goldman,G.H., Carvalho,A.E., Matsukuma,A., Baia,G.S., Simpson,D.H.,  
Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare  
M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and  
Simpson,A.J.  
Shotgun sequencing of the human transcriptome with ORF expressed  
sequence tags  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
20202663  
Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=PM3at2-PM3-HT0909-  
191100-028-f09et3=2000-11-19&t4=1)  
Seq primer: Puc 18 forward  
High quality sequence start: 27  
High quality sequence stop: 173.  
High quality sequence Location/Qualifiers  
1. .174  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_lib="HT0909"  
/dev\_stage="Adult"

FEATURES  
source  
1. 174  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_lib="HT0909"  
/dev\_stage="Adult"



allows identification of the library of origin of a clone within the mixture. The subtracted library (UI-R-CO) was constructed as follows: PCR amplified cDNA inserts from a pool of UI-R-AI and UI-R-EI clones from which 3' ESTs had been derived was used as a driver in a hybridization with the pooled UI-R-AI and UI-R-EI library in the form of single-stranded circles. The remaining single-stranded circles (subtracted library) was purified by hydroxyapatite column chromatography, converted to double-stranded circles and electroporated into DH10B bacteria (Life Technologies) to generate the UI-R-CO library. This procedure has been previously described (Bonaldi, Lennon and Soares, Genome Research 6: 791-806, 1996)."

BASE COUNT 40 a 31 c 47 g 59 t  
 ORIGIN

Query Match 100.0%; Score 12; DB 9; Length 177;  
 Best Local Similarity 100.0%; Pred. No. 3.6e+04;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
 |||||

Db 85 GTCGTGCAGGCA 96

RESULT 4  
 BE150969

LOCUS BE150969 180 bp mRNA linear EST 21-JUN-2000  
 DEFINITION RC4-HT0276-100300-015-f11 HT0276 Homo sapiens cDNA, mRNA sequence.  
 ACCESSION BE150969  
 VERSION BE150969.1 GI:8613690  
 KEYWORDS EST.  
 SOURCE human.

ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 180)  
 Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R., Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F., Goldman, G.H., Carvalho, A.F., Matsukuma, A., Baia, G.S., Simpson, D.H., Brunstein, A., deOliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.

TITLE Shotgun sequencing of the human transcriptome with ORF expressed sequence tags  
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
 MEDLINE 20202663  
 COMMENT

Contact: Simpson A.J.G.  
 Laboratory of Cancer Genetics  
 Ludwig Institute for Cancer Research  
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil  
 Tel: +55-11-2704922  
 Fax: +55-11-2707001  
 Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL  
 (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=at2=RC4-HT0276-100300-015-f11&t3=2000-03-10&t4=1)  
 Seq primer: puc 18 forward  
 High quality sequence start: 2  
 High quality sequence stop: 180.  
 Location/Qualifiers

FEATURES  
 source  
 1..180  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone\_lib="HT0276"  
 /dev\_stage="Adult"  
 /note="Organ: head\_neck; Vector: puc18; Site\_1: SmaI; Site\_2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research)

profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

BASE COUNT 31 a 47 c 51 g 51 t  
 ORIGIN

Query Match 100.0%; Score 12; DB 10; Length 180;  
 Best Local Similarity 100.0%; Pred. No. 3.6e+04;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
 |||||

Db 47 GTCGTGCAGGCA 58

RESULT 5  
 BI024277/c

LOCUS BI024277 192 bp mRNA linear EST 14-JUN-2001  
 DEFINITION CM3-WT0293-260101-691-h12 MT0293 Homo sapiens cDNA, mRNA sequence.  
 ACCESSION BI024277  
 VERSION BI024277.1 GI:14430907  
 KEYWORDS EST.  
 SOURCE human.

ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 192)  
 Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R., Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F., Goldman, G.H., Carvalho, A.F., Matsukuma, A., Baia, G.S., Simpson, D.H., Brunstein, A., deOliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.

TITLE Shotgun sequencing of the human transcriptome with ORF expressed sequence tags  
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
 MEDLINE 20202663  
 COMMENT

Contact: Simpson A.J.G.  
 Laboratory of Cancer Genetics  
 Ludwig Institute for Cancer Research  
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil  
 Tel: +55-11-2704922  
 Fax: +55-11-2707001  
 Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL  
 (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM3&t2=CM3-MT0293-260101-691-h12&t3=2001-01-26&t4=1)  
 Seq primer: puc 18 forward  
 High quality sequence start: 90  
 High quality sequence stop: 190.  
 Location/Qualifiers

FEATURES  
 source  
 1..192  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone\_lib="MT0293"  
 /dev\_stage="Adult"  
 /note="Organ: marrow; Vector: puc18; Site\_1: SmaI; Site\_2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

BASE COUNT 52 a 44 c 58 g 38 t  
 ORIGIN

Query Match 100.0%; Score 12; DB 13; Length 192;  
 Best Local Similarity 100.0%; Pred. No. 3.7e+04;  
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12

```

Db      81  GTCTGTCAGGCA 70
|||||
RESULT 6
BI034110/c
LOCUS      212 bp      mRNA      linear      EST 14-JUN-2001
DEFINITION PM2-NN0165-160301-007-e10 NN0165 Homo sapiens cDNA, mRNA sequence.
ACCESSION BI034110
VERSION    BI034110.1  GI:14440736
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare
,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.
Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?t1=PM2&t2=PM2-NN0165-
160301-007-e10&t3=2001-03-16&t4=1)
Seq primer: puc 18 forward
High quality sequence stop: 212.
Location/Qualifiers
1. .212
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="NN0165"
/dev_stage="Adult"
/note="Organ: nervous_normal; Vector: puc18; Site_1: SmaI;
Site_2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."
BASE COUNT 50 a 72 c 50 g 40 t
ORIGIN
Query Match 100.0%; Score 12; DB 13; Length 212;
Best Local Similarity 100.0%; Pred. No. 3.7e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1  GTCTGTCAGGCA 12
|||||
Db 172 GTCTGTCAGGCA 161
|||||
RESULT 7
AA815339
LOCUS      221 bp      mRNA      linear      EST 31-DEC-1998
DEFINITION ai62g07.s1 Soares_testis.NHT Homo sapiens cDNA clone 1375452 3'
similar to contains MER7.t3 MER7 repetitive element ;, mRNA
sequence.
ACCESSION AA815339
VERSION    AA815339.1  GI:2884935
|||||
Db      81  GTCTGTCAGGCA 70
|||||
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare
,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.
Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?t1=PM2&t2=PM2-NN0165-
160301-007-e10&t3=2001-03-16&t4=1)
Seq primer: puc 18 forward
High quality sequence stop: 212.
Location/Qualifiers
1. .221
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="1375452"
/clone_lib="Soares_testis_NHT"
/sex="male"
/lab_host="DH10B"
/note="Vector: pT73D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was prepared from mRNA obtained from Clontech Laboratories
, Inc., and primed with a Not I - oligo(dT) primer [5',
TGTTACCACTGAGCTGGAGCGGCCCAATTTTTTTTTTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization to Cot5, and was
constructed by Bento Soares and M. Fatima Bonaldo. "
BASE COUNT 57 a 47 c 52 g 65 t
ORIGIN
Query Match 100.0%; Score 12; DB 9; Length 221;
Best Local Similarity 100.0%; Pred. No. 3.7e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1  GTCTGTCAGGCA 12
|||||
Db 32  GTCTGTCAGGCA 43
|||||
RESULT 8
BF760820/c
LOCUS      239 bp      mRNA      linear      EST 12-JAN-2001
DEFINITION RC4-CT0109-311200-022-a05 CT0109 Homo sapiens cDNA, mRNA sequence.
ACCESSION BF760820
VERSION    BF760820.1  GI:12108720
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare
,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.
Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

```

MEDLINE  
COMMENT

20202663  
Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=RC4&t2=RC4-CT0109-  
311200-022-a05&t3=2000-12-31&t4=1)  
311200-022-a05&t3=2000-12-31&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 65  
High quality sequence stop: 239.

## FEATURES

Source

1..239  
Location/Qualifiers  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_lib="CT0109"  
/dev\_stage="Adult"  
/note="Organ: colon; Vector: puc18; Site\_1: SmaI; Site\_2:  
SmaI; A mini-library was made by cloning products derived  
from ORESTES PCR (U.S. Letters Patent application No. 196  
/716 - Ludwig Institute for Cancer Research) profiles  
into the pUC 18 vector. Reverse transcription of tissue  
mRNA and cDNA amplification were performed under low  
stringency conditions."

BASE COUNT 58 a 58 c 55 g 58 t  
ORIGIN

Query Match 100.0%; Score 12; DB 12; Length 239;  
Best Local Similarity 100.0%; Pred. No. 3.8e+04;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GTCGTCAGGCA 12  
|||||  
Db 42 GTCGTCAGGCA 31

RESULT 9  
BH227331/C  
LOCUS BH227331 248 bp DNA linear GSS 08-NOV-2001  
DEFINITION 1006139B07.y1 1006 - RescueMu Grid G Zea mays genomic, DNA  
sequence.  
ACCESSION BH227331  
VERSION BH227331.1 GI:16827224  
KEYWORDS GSS.  
SOURCE Zea mays.  
ORGANISM Zea mays

REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC  
clade; Panicoideae; Andropogoneae; Zea.  
1 (bases 1 to 248)  
Walbot V.  
Maize genomic sequences found using engineered RescueMu transposon  
Contact: Walbot V  
Department of Biological Sciences  
Stanford University  
855 California Ave, Palo Alto, CA 94304, USA  
Tel: 650 723 2227  
Fax: 650 725 8221  
Email: walbot@stanford.edu  
Possible ligation site so sequence was trimmed. Post-ligation  
sequence submitted separately.  
Plate: 1006139 row: 4  
Class: transposon-tagged.  
Location/Qualifiers  
1..248  
/organism="Zea mays"  
/cultivar="mixed background W23/A188/B73"

FEATURES  
Source

/db\_xref="taxon:4577"  
/clone\_lib="1006 - RescueMu Grid G"  
/tissue\_type="leaf"  
/dev\_stage="adult"  
/lab\_host="DH10B"  
/note="Organ: leaf; Vector: RescueMu (engineered from  
pBlueScript backbone); Site\_1: BamHI; Site\_2: BglII;  
RescueMu is a 4.9 kb, modified maize Mu transposon  
designed to allow plasmid rescue from total genomic DNA.  
Mu elements insert preferentially into transcription  
units. For more information on RescueMu, go to the web  
site 'www.zmdb.tastate.edu' and follow the links for  
'RescueMu.' Grid G was grown at Stanford in 2000. DNA was  
extracted from leaf punches, double digested using BamHI  
and BglII, and ligated to form circular plasmids. DH10B  
cells were transformed and then screened on LB plates with  
ampicillin." 40 a 80 c 89 g 39 t  
ORIGIN

Query Match 100.0%; Score 12; DB 17; Length 248;  
Best Local Similarity 100.0%; Pred. No. 3.8e+04;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GTCGTCAGGCA 12  
|||||  
Db 62 GTCGTCAGGCA 51

RESULT 10  
BM145815/C  
LOCUS BM145815 253 bp mRNA linear EST 30-NOV-2001  
DEFINITION TCAAPID7858 Pediatric acute myelogenous leukemia cell (FAB M1)  
Baylor-HGSC project=TCAA Homo sapiens cDNA clone TCAAP7858, mRNA  
sequence.  
ACCESSION BM145815  
VERSION BM145815.1 GI:17164180  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 253)  
AUTHORS Wei Y., Tsang Y.T.M., Mei G., Ku J.M., Ali-Osman F.R. Jr.,  
Gundaratne P.H., Muzny D., Bouck J., Gibbs R.A. and Margolin J.F.  
TITLE Pediatric Leukemia cDNA Sequencing Project (2001)  
JOURNAL Unpublished (2001)  
COMMENT Contact: Dr. Judith F. Margolin  
Texas Children's Cancer Center and Human Genome Sequencing Center  
at Baylor College of Medicine  
1102 Bates, MC3-3320 Houston, TX 77030, USA  
Tel: 832-824-4536  
Fax: 832-825-4038  
Email: clones@tccc.org  
Seq primer: M13 primer.

FEATURES  
Source

1..253  
Location/Qualifiers  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="TCAAP7858"  
/clone\_lib="Pediatric acute myelogenous leukemia cell (FAB  
M1) Baylor-HGSC project=TCAA"  
/sex="male"  
/tissue\_type="leukopheresis"  
/cell\_type="myeloid cell"  
/dev\_stage="pediatric 6 years"  
/lab\_host="DH10B"  
/note="Vector: lambda pSB; Site\_1: BamHI; Site\_2: EcoRI;  
First strand cDNA was primed with an anchored  
XhoI-oligo(dT) primer [5'GGAGGACTCGAGCGCCGAGGAG(T)VN  
3'; V=A,C,G; N=A,C,G,T] and then dg tailed. Second strand  
was primed with a BamHI-dC primer  
[5'AGAGAGCTCGGATCGCGCGCCCAATAATAAT(C) 3'].

Mon Jan 6 15:20:20 2003

Double-stranded cDNA was then digested with BamHI and XhoI and directionally cloned into the BamHI and SalI sites of lambda PSB vector. Library was constructed by one round of normalization. Library was constructed by Wei Yu at RIKEN of Japan (Carninci P, Westover A, Nishiyama Y, Ohsumi T, Itoh M, Nagaoka S, Sasaki N, Okazaki Y, Muramatsu M, Schneider C, Hayashizaki Y, High efficiency selection of full-length cDNA by improved biotinylated cap trapper., DNA Res 4: 1, 61-6, Feb 28, 1997)"

34 a 65 c 85 g 67 t 2 others

Query Match 100.0%; Score 12; DB 13; Length 253;  
Best Local Similarity 100.0%; Pred. No. 3.8e+04;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12

|||||

Db 218 GTCGTGCAGGCA 207

RESULT 11  
BF256224/c  
LOCUS  
DEFINITION  
HVSMEf0009D01f Hordeum vulgare seedling root EST library HVCdNA0007 (Etolated and unstressed) Hordeum vulgare cDNA clone  
HVSMEf0009D01f, mRNA sequence.

ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Hordeum vulgare.  
Hordeum vulgare.  
EST.  
Hordeum vulgare.

REFERENCE  
AUTHORS  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae; Triticeae; Hordeum.

TITLE  
JOURNAL  
COMMENT  
1 (bases 1 to 256)  
Wing R., Close T.J., Kleinhoofs A., Wise R., Begum D., Frisch D., Yu Y., Henry D., Palmer M., Rambo T., Simmons J., Choi D.W., Fenton R.D., Oates R. and Main D.  
Development of a genetically and physically anchored EST resource for barley genomics: Morex unstressed seedling root cDNA library  
Unpublished (2001)  
On Nov 16, 2000 this sequence version replaced gi:11185337.  
Contact: Wing RA

Location/Qualifiers  
1. .256  
/organism="Hordeum vulgare"  
/cultivar="Morex"  
/db\_xref="taxon:4513"  
/clone="HVSMEf0009D01f"  
HVCdNA0007 (Etolated and unstressed)  
/tissue\_type="Seedling root"  
/lab\_host="TJC121"  
/note="Vector: lambdaZAP; Site 1: EcoRI; Site 2: XhoI;  
Seeds were surface sterilized then germinated under axenic conditions in the dark at room temperature on filter paper with water, nystatin and ceftaxime in covered crystallization dishes. Five-day old seedling roots were then harvested, total RNA was prepared, poly(A) RNA was purified, one primary unamplified cDNA library was made, and 1 million pfu were in vivo excised to give pBluescript SK(-) cDNA phagemids. These steps were performed in the TJ

FEATURES  
source

MG1:1422433  
Possible reversed clone: polyT not found  
Seq primer: -400P from Gibco.  
Location/Qualifiers  
1. .261  
/organism="Mus musculus"  
/strain="CZECH II"  
/db\_xref="taxon:10090"  
/clone="IMAGE:3661665"  
/clone\_lib="NCI\_CGAP-Lu30"  
/tissue\_type="tumor, metastatic to mammary"  
/lab\_host="DH10B"  
/note="Organ: lung; Vector: pCMV-SPORT6; Site 1: NotI;  
Site 2: SalI; transgenic model WNT-1, expression driven by MMTV-LTR enhancer; Cloned unidirectionally. Primer: Oligo dT. Library constructed by Life Technologies.  
Investigator providing samples: Gilbert Smith, NIH"

Close laboratory at the University of California, Riverside (Choi, Close, Fenton). Phagemids were plated and picked at the Clemson University Genomics Institute (CUGI) (Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA preparations, DNA sequencing and sequence analysis were performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main). The sequence has been trimmed to remove vector sequence and contains a minimum of 100 bases of phred value 20 or above. For more details on library preparation and sequence analysis see <http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> Also see Close TJ, Wing R, Kleinhoofs A, Wise R (2001) Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html>.)

BASE COUNT 57 a 84 c 73 g 42 t  
ORIGIN  
Query Match 100.0%; Score 12; DB 12; Length 256;  
Best Local Similarity 100.0%; Pred. No. 3.8e+04;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12

|||||

Db 221 GTCGTGCAGGCA 210

RESULT 12  
BF146813/c  
LOCUS  
DEFINITION  
BF146813 261 bp mRNA linear EST 29-DEC-2000  
uy36f05.x1 NCI\_CGAP-Lu30 Mus musculus cDNA clone IMAGE:3661665 3' similar to SW:SMB2\_MOUSE P40694 DNA-BINDING PROTEIN SMUBP-2 ; mRNA sequence.

ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
BF146813.1 GI:11028208  
EST.  
house mouse.  
Mus musculus

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT  
1 (bases 1 to 261)  
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
Unpublished (1997)  
Contact: Robert Strausberg, Ph.D.  
Email: cgapbs-r@mail.nih.gov  
Tissue Procurement: Gilbert Smith, Ph.D.  
cDNA Library Preparation: Life Technologies, Inc.  
DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)  
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: [image.llnl.gov/image/html/iresources.shtml](http://image.llnl.gov/image/html/iresources.shtml)

MG1:1422433  
Possible reversed clone: polyT not found  
Seq primer: -400P from Gibco.  
Location/Qualifiers  
1. .261  
/organism="Mus musculus"  
/strain="CZECH II"  
/db\_xref="taxon:10090"  
/clone="IMAGE:3661665"  
/clone\_lib="NCI\_CGAP-Lu30"  
/tissue\_type="tumor, metastatic to mammary"  
/lab\_host="DH10B"  
/note="Organ: lung; Vector: pCMV-SPORT6; Site 1: NotI;  
Site 2: SalI; transgenic model WNT-1, expression driven by MMTV-LTR enhancer; Cloned unidirectionally. Primer: Oligo dT. Library constructed by Life Technologies.  
Investigator providing samples: Gilbert Smith, NIH"



Mon Jan 6 15:20:20 2003

priming with dT-tailed vector. The dT-tailed vector was adjusted to have about 60nt. The cDNA vector was circularized with E. coli DNA ligase after digestion of EcoRI which site is also included in vector. An RNA strand converted to a DNA strand by Okayama-Berg method. The obtained cDNA vectors were used for transformation of competent cells E. coli Top10F<sup>+</sup> by electroporation method. The cDNA libraries constructed by this method are full-length enriched cDNA library."

BASE COUNT 39 a 96 c 80 g 77 t

ORIGIN

Query Match 100.0%; Score 12; DB 14; Length 292;

Best Local Similarity 100.0%; Pred. No. 3.9e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12

|||||

Db 32 GTCGTGCAGGCA 43

RESULT 15

AT004345/C

LOCUS

DEFINITION

AT004345

sequence.

AT004345

VERSION

AT004345

KEYWORDS

EST.

SOURCE

Oyster mushroom.

ORGANISM

Pleurotus ostreatus

Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes;

Agaricales; Pleurotaceae; Pleurotus.

1 (bases 1 to 296)

Lee, S.H., Kim, B.G., Kim, K.J., Lee, J.S., Yun, D.W., Hahn, J.H., Kim,

G.H., Lee, K.H., Suh, D.S., Kwon, S.T., Lee, C.S. and Yoo, Y.B.

Comparative Analysis of Sequences Expressed during the

Liquid-Cultured Mycelia and Fruit Body Stages of Pleurotus

ostreatus

Fungal Genet. Biol. 35 (2), 115-134 (2002)

21838665

Contact: Beom-Gi Kim

Division of applied microbiology

Institute of Agricultural Science and Technology(NIAST)

249 Seodundong Kwonseonku, Suwon 441707, Korea

Tel: 82-331-290-0347

Fax: 82-331-290-0399

Email: bskimyes@da.go.kr

GeneNuri No. KSL04602

Submitted through BRIC(Biological Research Information Center) of

Korea

URL: http://bric.postech.ac.kr/.

Location/Qualifiers

1. .296

/organism="Pleurotus ostreatus"

/cultivar="AST 2029"

/db\_xref="taxon:5322"

/clone="1893LM"

/clone\_lib="POSLM01"

/dev\_stage="Shaking liquid cultured mycelia"

/lab\_host="E.coli"

/note="Vector: lambda Uni-ZAP XR; Site.1: EcoRI; Site.2:

XhoI; average insert size:1000 bp;Initial pfu:5 X 10<sup>7</sup>

Library information:Isolation of total RNA from the

mycelia incubated in shaking liquid MCM media at 30 deg C"

BASE COUNT 97 a 80 c 60 g 57 t 2 others

ORIGIN

Query Match 100.0%; Score 12; DB 9; Length 296;

Best Local Similarity 100.0%; Pred. No. 3.9e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12

Db 273 GTCGTGCAGGCA 262

Search completed: January 4, 2003, 01:04:23

Job time : 164.558 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:53:16 ; Search time 4.05047 Seconds  
(without alignments)  
908.566 Million cell updates/sec

Title: US-09-787-562-10  
Perfect score: 12  
Sequence: 1 gtcgtgcaggca 12

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 15338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued\_Patents\_NA.\*  
1: /cgn2\_6/ptodata/2/ina/5A\_COMB.seq.\*  
2: /cgn2\_6/ptodata/2/ina/5B\_COMB.seq.\*  
3: /cgn2\_6/ptodata/2/ina/6A\_COMB.seq.\*  
4: /cgn2\_6/ptodata/2/ina/6B\_COMB.seq.\*  
5: /cgn2\_6/ptodata/2/ina/PTUS\_COMB.seq.\*  
6: /cgn2\_6/ptodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	12	100.0	35	1	US-08-360-841-4
C 2	12	100.0	36	1	US-08-360-841-11
C 3	12	100.0	174	1	US-08-360-841-1
C 4	12	100.0	953	4	US-08-438-745-3
C 5	12	100.0	953	4	US-08-438-745-5
C 6	12	100.0	953	4	US-08-219-019-3
C 7	12	100.0	953	4	US-09-219-019-5
C 8	12	100.0	953	5	PCT-US94-05669A-3
C 9	12	100.0	953	5	PCT-US94-05669A-5
C 10	12	100.0	2087	4	US-09-097-199-83
C 11	12	100.0	7859	2	US-07-854-956B-4
C 12	12	100.0	7859	2	US-08-450-905B-15
C 13	12	100.0	7859	3	US-07-982-759F-15
C 14	11	91.7	30	4	US-08-327-984A-13
C 15	11	91.7	30	4	US-09-327-984A-14
C 16	11	91.7	41	4	US-09-327-984A-28
C 17	11	91.7	49	4	US-09-327-984A-15
C 18	11	91.7	49	4	US-09-327-984A-16
C 19	11	91.7	49	4	US-09-327-984A-17
C 20	11	91.7	49	4	US-09-327-984A-18
C 21	11	91.7	49	4	US-09-327-984A-19
C 22	11	91.7	282	2	US-08-105-989-10
C 23	11	91.7	282	3	US-09-138-922-10
C 24	11	91.7	416	4	US-09-319-056B-1
C 25	11	91.7	416	4	US-09-319-056B-3
C 26	11	91.7	614	2	US-08-729-103-2
C 27	11	91.7	652	4	US-08-998-416-962

28	11	91.7	861	4	US-08-998-416-299	Sequence 299, Appl
C 29	11	91.7	957	2	US-08-544-822-2	Sequence 2, Appli
C 30	11	91.7	957	3	US-09-070-964-2	Sequence 2, Appli
31	11	91.7	1080	3	US-09-188-930-9	Sequence 9, Appli
32	11	91.7	1080	4	US-09-125-642C-3	Sequence 3, Appli
C 33	11	91.7	1114	2	US-08-468-413-1	Sequence 1, Appli
C 34	11	91.7	1114	3	US-09-162-508-1	Sequence 1, Appli
C 35	11	91.7	1114	5	PCT-US95-07169-1	Sequence 1, Appli
C 36	11	91.7	1193	1	US-07-956-697B-4	Sequence 4, Appli
C 37	11	91.7	1193	1	US-08-263-098-4	Sequence 4, Appli
38	11	91.7	1193	4	US-09-541-941B-27	Sequence 27, Appl
C 39	11	91.7	1246	1	US-08-446-777-3	Sequence 3, Appli
C 40	11	91.7	1249	4	US-09-333-208-1	Sequence 1, Appli
C 41	11	91.7	1249	4	US-09-333-254-1	Sequence 1, Appli
C 42	11	91.7	1249	4	US-09-183-270-1	Sequence 1, Appli
43	11	91.7	1280	3	US-09-188-930-246	Sequence 246, App
44	11	91.7	1445	1	US-08-324-533-1	Sequence 1, Appli
45	11	91.7	1501	3	US-08-993-359-21	Sequence 21, Appl

## ALIGNMENTS

RESULT 1  
US-08-360-841-4/c  
; Sequence 4, Application US/08360841  
; Patent No. 5652120  
; GENERAL INFORMATION:  
; APPLICANT: PARK, Seung Kook  
; APPLICANT: LEE, Kang Moon  
; APPLICANT: NHO, Kyoo Seung  
; APPLICANT: KOH, Yeo Wook  
; APPLICANT: KWON, Chang Hyuk  
; APPLICANT: CHUNG, Ju Young  
; APPLICANT: JEE, Young Su  
; APPLICANT: YU, Young Hyo  
; TITLE OF INVENTION: A No. 5652120e1 Gene Coding Human Epidermal  
; TITLE OF INVENTION: Growth Factor and Process for Preparing the Same  
; NUMBER OF SEQUENCES: 20  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: darby & darby PC  
; STREET: 805 Third Avenue  
; CITY: New York  
; STATE: New York  
; COUNTRY: US  
; ZIP: 10022  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/360,841  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Ludwig, S. Peter  
; REGISTRATION NUMBER: 25,351  
; REFERENCE/DOCKET NUMBER: 0136/0A760  
; TELEPHONE: 212-527-7700  
; TELEFAX: 212-753-6237  
; TELEX: 236687  
; INFORMATION FOR SEQ ID NO: 4:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 35 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: cdna to mRNA  
; ORIGINAL SOURCE:  
; ORGANISM: Homo sapiens  
; IMMEDIATE SOURCE:

; CLONE: C2 PRIMER  
US-08-360-841-4

Query Match 100.0%; Score 12; DB 1; Length 35;  
Best Local Similarity 100.0%; Pred. No. 2.3e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12  
|||||  
Db 24 GTCGTGCAGGCA 13

## RESULT 2

US-08-360-841-11  
; Sequence 11, Application US/08360841  
; Patent No. 5652120

## ; GENERAL INFORMATION:

; APPLICANT: PARK, Seung Kook  
; APPLICANT: LEE, Kang Moon  
; APPLICANT: NHO, Kyo Seung  
; APPLICANT: KOH, Yeo Wook  
; APPLICANT: KWON, Chang Hyuk  
; APPLICANT: CHUNG, Ju Young  
; APPLICANT: JEE, Young Su

; APPLICANT: Yu, Young Hyo

; TITLE OF INVENTION: A No. 5652120el Gene Coding Human Epidermal

; GROWTH FACTOR AND PROCESS FOR PREPARING THE SAME

; NUMBER OF SEQUENCES: 20

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Darby & Darby PC

; STREET: 805 Third Avenue

; CITY: New York

; STATE: New York

; COUNTRY: US

; ZIP: 10022

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/360,841

; FILING DATE:

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Ludwig, S. Peter

; REGISTRATION NUMBER: 25,351

; REFERENCE/DOCKET NUMBER: 0136/OA760

; TELEPHONE: 212-527-7700

; TELEFAX: 212-753-6237

; TELEX: 236687

; INFORMATION FOR SEQ ID NO: 11:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 36 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: cDNA to mRNA

; ORIGINAL SOURCE:

; ORGANISM: Homo sapiens

; IMMEDIATE SOURCE:

; CLONE: N4 PRIMER

US-08-360-841-11

Query Match 100.0%; Score 12; DB 1; Length 36;  
Best Local Similarity 100.0%; Pred. No. 2.3e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12  
|||||  
Db 21 GTCGTGCAGGCA 32

## RESULT 3

US-08-360-841-1/c

; Sequence 1, Application US/08360841

; Patent No. 5652120

; GENERAL INFORMATION:

; APPLICANT: PARK, Seung Kook

; APPLICANT: LEE, Kang Moon

; APPLICANT: NHO, Kyo Seung

; APPLICANT: KOH, Yeo Wook

; APPLICANT: KWON, Chang Hyuk

; APPLICANT: CHUNG, Ju Young

; APPLICANT: JEE, Young Su

; APPLICANT: Yu, Young Hyo

; TITLE OF INVENTION: A No. 5652120el Gene Coding Human Epidermal

; GROWTH FACTOR AND PROCESS FOR PREPARING THE SAME

; NUMBER OF SEQUENCES: 20

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Darby & Darby PC

; STREET: 805 Third Avenue

; CITY: New York

; STATE: New York

; COUNTRY: US

; ZIP: 10022

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/360,841

; FILING DATE:

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Ludwig, S. Peter

; REGISTRATION NUMBER: 25,351

; REFERENCE/DOCKET NUMBER: 0136/OA760

; TELEPHONE: 212-527-7700

; TELEFAX: 212-753-6237

; TELEX: 236687

; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 174 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: cDNA to mRNA

; ORIGINAL SOURCE:

; ORGANISM: Homo sapiens

; IMMEDIATE SOURCE:

; CLONE: EGF-nt seq

US-08-360-841-1

Query Match 100.0%; Score 12; DB 1; Length 174;

Best Local Similarity 100.0%; Pred. No. 2.2e+02;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12

|||||

Db 54 GTCGTGCAGGCA 43

## RESULT 4

US-08-438-745-3/c

; Sequence 3, Application US/08438745

; Patent No. 6248715

; GENERAL INFORMATION:

; APPLICANT: Rosenberg, Steven

; APPLICANT: Stratton-Thomas, Jennifer

; TITLE OF INVENTION: Expression of Urokinase Plasminogen

; ACTIVATOR INHIBITORS

; NUMBER OF SEQUENCES: 22



;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: Chiron Corporation  
;; STREET: 4560 Horton Street  
;; CITY: Emeryville  
;; STATE: CA  
;; COUNTRY: USA  
;; ZIP: 94608  
;; COMPUTER READABLE FORM:  
;; MEDIUM TYPE: Floppy disk  
;; COMPUTER: IBM PC compatible  
;; OPERATING SYSTEM: PC-DOS/MS-DOS  
;; SOFTWARE: PatentIn Release #1.0, Version #1.25  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: US/08/438,745  
;; FILING DATE: 10-MAY-1995  
;; CLASSIFICATION: 514  
;; PRIOR APPLICATION NUMBER: US 08/070,153  
;; FILING DATE: 01-JUN-1993  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Green, Grant D.  
;; REGISTRATION NUMBER: 31,259  
;; REFERENCE/DOCKET NUMBER: 0939.001  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: 510-601-2706  
;; TELEFAX: 510-655-3542  
;; INFORMATION FOR SEQ ID NO: 3:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 953 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: double  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: cdna  
;; HYPOTHETICAL: NO  
;; IMMEDIATE SOURCE:  
;; CLONE: M1Flag-EGF-pIII fusion  
;; FEATURE:  
;; NAME/KEY: CDS  
;; LOCATION: 25..903  
;; US-08-438-745-3

Query Match 100.0%; Score 12; DB 4; Length 953;  
Best Local Similarity 100.0%; Pred. No. 2.2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
|||||  
Db 165 GTCGTGCAGGCA 154

RESULT 5  
US-08-438-745-5/c  
; Sequence 5, Application US/08438745  
; Patent No. 6248715  
; GENERAL INFORMATION:  
; APPLICANT: Rosenberg, Steven  
; TITLE OF INVENTION: Expression of Urokinase Plasminogen  
; TITLE OF INVENTION: Activator Inhibitors  
; NUMBER OF SEQUENCES: 22  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Chiron Corporation  
; STREET: 4560 Horton Street  
; CITY: Emeryville  
; STATE: CA  
; COUNTRY: USA  
; ZIP: 94608  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:

;; APPLICATION NUMBER: US/08/438,745  
;; FILING DATE: 10-MAY-1995  
;; CLASSIFICATION: 514  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 08/070,153  
;; FILING DATE: 01-JUN-1993  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Green, Grant D.  
;; REGISTRATION NUMBER: 31,259  
;; REFERENCE/DOCKET NUMBER: 0939.001  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: 510-601-2706  
;; TELEFAX: 510-655-3542  
;; INFORMATION FOR SEQ ID NO: 5:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 953 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: double  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: cdna  
;; HYPOTHETICAL: NO  
;; IMMEDIATE SOURCE:  
;; CLONE: M1Flag-EGF-pIII fusion  
;; FEATURE:  
;; NAME/KEY: CDS  
;; LOCATION: 25..903  
;; US-08-438-745-5

Query Match 100.0%; Score 12; DB 4; Length 953;  
Best Local Similarity 100.0%; Pred. No. 2.2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
|||||  
Db 165 GTCGTGCAGGCA 154

RESULT 6  
US-09-219-019-3/c  
; Sequence 3, Application US/09219019  
; Patent No. 6268341  
; GENERAL INFORMATION:  
; APPLICANT: ROSENBERG, STEVEN  
; APPLICANT: STRATTON-THOMAS, JENNIFER R.  
; TITLE OF INVENTION: EXPRESSION OF UROKINASE PLASMINOGEN ACTIVATOR  
; TITLE OF INVENTION: INHIBITORS  
; FILE REFERENCE: 23533-0005  
; CURRENT APPLICATION NUMBER: US/09/219,019  
; CURRENT FILING DATE: 1998-12-23  
; PRIOR APPLICATION NUMBER: 08/438,263  
; PRIOR FILING DATE: 1995-05-10  
; PRIOR APPLICATION NUMBER: 08/280,288  
; PRIOR FILING DATE: 1994-07-26  
; PRIOR APPLICATION NUMBER: 08/070,153  
; PRIOR FILING DATE: 1993-06-01  
; NUMBER OF SEQ ID NOS: 22  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 3  
; LENGTH: 953  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: (25)..(903)  
; US-09-219-019-3

Query Match 100.0%; Score 12; DB 4; Length 953;  
Best Local Similarity 100.0%; Pred. No. 2.2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
|||||  
Db 165 GTCGTGCAGGCA 154

us-09-787-562-10.rni

Mon Jan 6 15:20:18 2003

```

; LENGTH: 953 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cdna
; HYPOTHETICAL: NO
; IMMEDIATE SOURCE:
; CLONE: M1flag-EGF-pIII fusion
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 25..903
PCT-US94-05669A-3

Query Match 100.0%; Score 12; DB 5; Length 953;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12
Db 165 GTCGTGCAGGCA 154

RESULT 9
PCT-US94-05669A-5/c
; Sequence 5, Application PC/TUS9405669A
; GENERAL INFORMATION:
; APPLICANT: Chiron Corporation
; TITLE OF INVENTION: Expression of Urokinase Plasminogen
; TITLE OF INVENTION: Activator Inhibitors
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/05669A
; FILING DATE: 19-MAY-1994
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Green, Grant D.
; REGISTRATION NUMBER: 31,259
; REFERENCE/DOCKET NUMBER: 0939,100
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 510-601-2706
; TELEFAX: 510-655-3542
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 953 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cdna
; HYPOTHETICAL: NO
; IMMEDIATE SOURCE:
; CLONE: M1flag-EGF-pIII fusion
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 25..903
PCT-US94-05669A-5

Query Match 100.0%; Score 12; DB 5; Length 953;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12

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```

; LENGTH: 953
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (25)..(903)
US-09-219-019-5

Query Match 100.0%; Score 12; DB 4; Length 953;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTCGTGCAGGCA 12
Db 165 GTCGTGCAGGCA 154

RESULT 8
PCT-US94-05669A-3/c
; Sequence 3, Application PC/TUS9405669A
; GENERAL INFORMATION:
; APPLICANT: Chiron Corporation
; TITLE OF INVENTION: Expression of Urokinase Plasminogen
; TITLE OF INVENTION: Activator Inhibitors
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/05669A
; FILING DATE: 19-MAY-1994
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Green, Grant D.
; REGISTRATION NUMBER: 31,259
; REFERENCE/DOCKET NUMBER: 0939,100
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 510-601-2706
; TELEFAX: 510-655-3542
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:

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Db 165 GTCGTGAGGCA 154  
|||||

## RESULT 10

US-09-097-199-83/c  
; Sequence 83, Application US/09097199  
; Patent No. 6218529  
; GENERAL INFORMATION:  
; APPLICANT: An, Gang  
; APPLICANT: O'Hara, S. Mark  
; APPLICANT: Ralph, David  
; APPLICANT: Veltri, Robert  
; TITLE OF INVENTION: BIOMARKERS AND TARGETS FOR DIAGNOSIS,  
; PROGNOSIS AND MANAGEMENT OF PROSTATE DISEASE  
; NUMBER OF SEQUENCES: 87  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Arnold, White & Durkee  
; STREET: P.O. Box 4433  
; CITY: Houston  
; STATE: Texas  
; COUNTRY: USA  
; ZIP: 77210  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/097,199  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/692,787  
; FILING DATE: 31-JUL-1996  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Nakashima, Richard A.  
; REGISTRATION NUMBER: P-42,023  
; REFERENCE/DOCKET NUMBER: UROC:018  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (512) 418-3000  
; TELEFAX: (512) 474-7577  
; INFORMATION FOR SEQ ID NO: 83:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2087 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 99..503  
; US-09-097-199-83

Query Match 100.0%; Score 12; DB 4; Length 2087;  
Best Local Similarity 100.0%; Pred. No. 2.2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGAGGCA 12  
|||||

Db 1967 GTCGTGAGGCA 1956

## RESULT 11

US-07-854-596B-4/c  
; Sequence 4, Application US/07854596B  
; Patent No. 5434073  
; GENERAL INFORMATION:  
; APPLICANT: Dawson, Keith M  
; APPLICANT: Hunter, Michael G  
; APPLICANT: Czaplowski, Lloyd G  
; TITLE OF INVENTION: Proteins and nucleic acids  
; NUMBER OF SEQUENCES: 73  
; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Dr. John J. McDonnell  
; STREET: Ten South Wacker Drive, Suite 3000  
; CITY: Chicago  
; STATE: IL  
; COUNTRY: USA  
; ZIP: 60606  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/07/854,596B  
; FILING DATE: 03-JUN-1992  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: McDonnell, John J  
; REGISTRATION NUMBER: 26,949  
; REFERENCE/DOCKET NUMBER: 92,337  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 312-715-1000  
; TELEFAX: 312-715-1234  
; TELEX: 910-221-5317  
; INFORMATION FOR SEQ ID NO: 4:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 7859 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: circular  
; MOLECULE TYPE: cDNA  
; FEATURE:  
; NAME/KEY: misc feature  
; LOCATION: 1..7859  
; OTHER INFORMATION: /note= "sequence of plasmid psw6"  
; US-07-854-596B-4

Query Match 100.0%; Score 12; DB 1; Length 7859;  
Best Local Similarity 100.0%; Pred. No. 2.2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGAGGCA 12  
|||||

Db 7739 GTCGTGAGGCA 7728

## RESULT 12

US-08-450-905B-15/c  
; Sequence 15, Application US/08450905B  
; Patent No. 5856301  
; GENERAL INFORMATION:  
; APPLICANT:  
; TITLE OF INVENTION: Stem Cell Inhibiting Proteins  
; NUMBER OF SEQUENCES: 178  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: HALE and DORR  
; STREET: 60 State Street  
; CITY: Boston  
; STATE: MA  
; ZIP: 02109  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/450,905B  
; FILING DATE: 26-MAR-1995  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 07/982,759  
; FILING DATE: 08-MAR-1993  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: GB 9127319.3  
; FILING DATE: 23-DEC-1991

```
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9221587.0
; FILING DATE: 14-OCT-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: BAKER, HOLLIE L.
; REGISTRATION NUMBER: 31,321
; REFERENCE/DOCKET NUMBER: 102.378.120DV-2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-526-6110
; TELEFAX: 617-526-5000
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7859 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA
US-08-450-905B-15

Query Match 100.0%; Score 12; DB 2; Length 7859;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
|||||
Db 7739 GTCGTGCAGGCA 7728

RESULT 13
US-07-982-759F-15/c
; Sequence 15, Application US/07982759F
; Patent No. 6057123
; GENERAL INFORMATION:
; APPLICANT: CRAIG, Stewart
; APPLICANT: GEORGE, Michael
; APPLICANT: EDWARDS, Richard Mark
; APPLICANT: CZAPLEWSKI, Lloyd George
; APPLICANT: GILBERT, Richard
; TITLE OF INVENTION: Stem Cell Inhibiting Proteins
; NUMBER OF SEQUENCES: 178
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HALE and DORR LLP
; STREET: 60 State Street
; CITY: Boston
; STATE: MA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/982.759F
; FILING DATE: 08-MAR-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9127319.3
; FILING DATE: 23-DEC-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9221587.0
; FILING DATE: 14-OCT-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: BAKER, HOLLIE L.
; REGISTRATION NUMBER: 31,321
; REFERENCE/DOCKET NUMBER: 102378.120
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-526-6000
; TELEFAX: 617-526-5000
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7859 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
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; MOLECULE TYPE: DNA
US-07-982-759F-15

Query Match 100.0%; Score 12; DB 3; Length 7859;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12
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Db 7739 GTCGTGCAGGCA 7728

RESULT 14
US-09-327-984A-13/c
; Sequence 13, Application US/09327984A
; Patent No. 6368594
; GENERAL INFORMATION:
; APPLICANT: Doetsch, Paul W.
; APPLICANT: Kaur, Balveen
; APPLICANT: Avery, Angela M.
; TITLE OF INVENTION: Broad Specificity DNA Damage Endonuclease
; FILE REFERENCE: 25-98
; CURRENT APPLICATION NUMBER: US/09/327,984A
; CURRENT FILING DATE: 1999-06-08
; PRIOR APPLICATION NUMBER: US 60/088,521
; PRIOR FILING DATE: 1998-06-08
; PRIOR APPLICATION NUMBER: US 60/134,752
; PRIOR FILING DATE: 1999-05-18
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 13
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence:double stranded
; OTHER INFORMATION: oligonucleotide containing cis-syn cyclobutane
; OTHER INFORMATION: pyrimidine dimer
; NAME/KEY: misc.feature
; LOCATION: (15)..(16)
; OTHER INFORMATION: At positions 15-16, the T-T is in the form of a
; OTHER INFORMATION: cis-syn cyclobutane pyrimidine dimer
US-09-327-984A-13

Query Match 91.7%; Score 11; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 8.5e+02;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCGTGCAGGCA 12
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Db 13 TCGTGCAGGCA 3

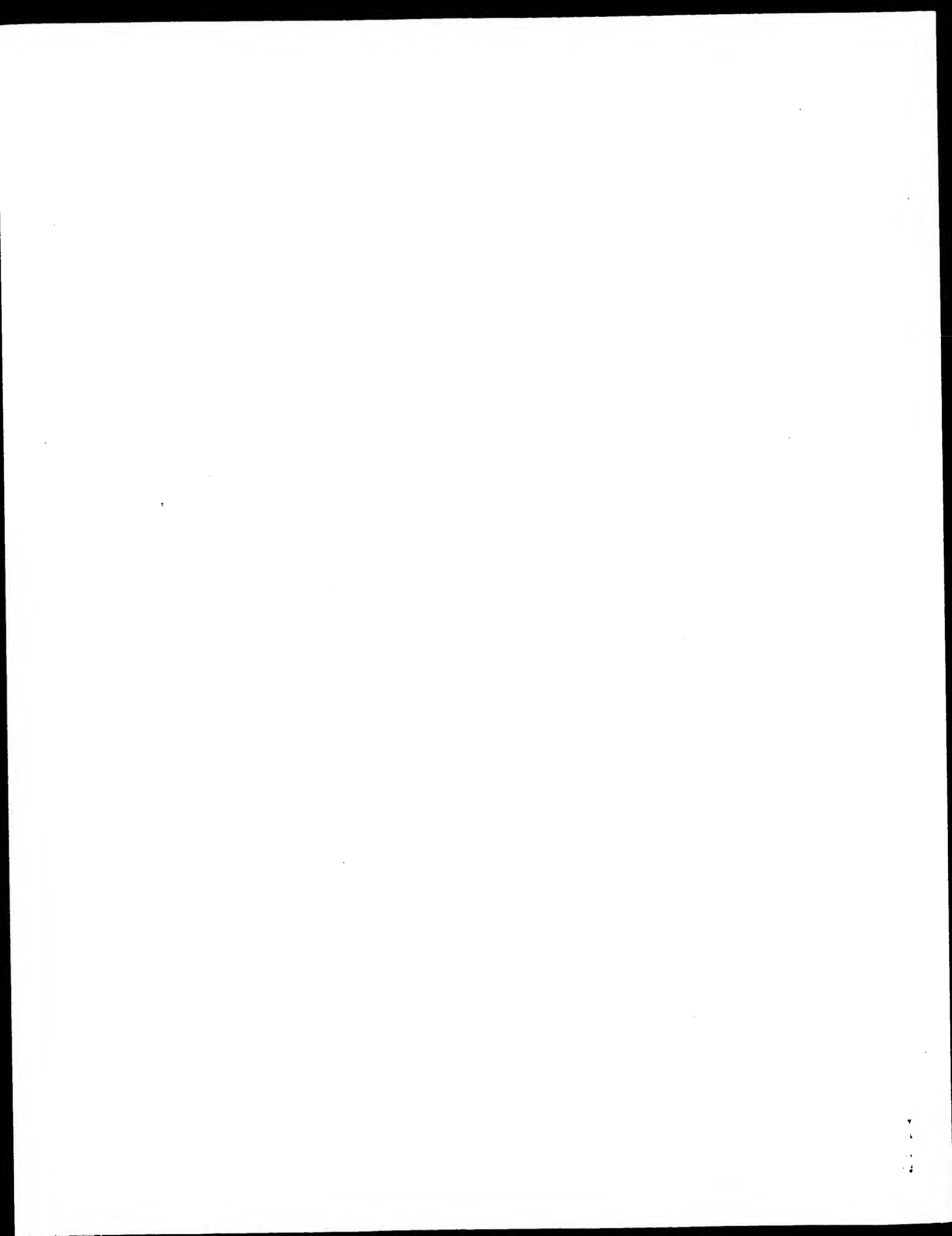
RESULT 15
US-09-327-984A-14/c
; Sequence 14, Application US/09327984A
; Patent No. 6368594
; GENERAL INFORMATION:
; APPLICANT: Doetsch, Paul W.
; APPLICANT: Kaur, Balveen
; APPLICANT: Avery, Angela M.
; TITLE OF INVENTION: Broad Specificity DNA Damage Endonuclease
; FILE REFERENCE: 25-98
; CURRENT APPLICATION NUMBER: US/09/327,984A
; CURRENT FILING DATE: 1999-06-08
; PRIOR APPLICATION NUMBER: US 60/088,521
; PRIOR FILING DATE: 1998-06-08
; PRIOR APPLICATION NUMBER: US 60/134,752
; PRIOR FILING DATE: 1999-05-18
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14
; LENGTH: 30
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; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence:undamaged
; OTHER INFORMATION: double stranded oligonucleotide
US-09-327-984A-14
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Query Match          91.7%; Score 11; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 8.5e+02;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 2 TCGTGCAGGCA 12
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Db 13 TCGTGCAGGCA 3
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Search completed: January 4, 2003, 00:10:07
Job time : 6.05047 secs
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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:41 ; Search time 4.05047 Seconds  
(without alignments)  
1281.345 Million cell updates/sec

Title: US-09-787-562-10

Perfect score: 12

Sequence: 1 gtcgtcaggca 12

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 381593 seqs, 216252194 residues

Total number of hits satisfying chosen parameters: 763186

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PublishedApplications.NA.\*

- 1: /cgn2\_6/ptodata/1/pubpna/US07\_PUBCOMB.seq.\*
- 2: /cgn2\_6/ptodata/1/pubpna/PTCT\_NEW\_PUB.seq.\*
- 3: /cgn2\_6/ptodata/1/pubpna/US06\_NEW\_PUB.seq.\*
- 4: /cgn2\_6/ptodata/1/pubpna/US06\_PUBCOMB.seq.\*
- 5: /cgn2\_6/ptodata/1/pubpna/US07\_NEW\_PUB.seq.\*
- 6: /cgn2\_6/ptodata/1/pubpna/PTCTUS\_PUBCOMB.seq.\*
- 7: /cgn2\_6/ptodata/1/pubpna/US08\_NEW\_PUB.seq.\*
- 8: /cgn2\_6/ptodata/1/pubpna/US08\_PUBCOMB.seq.\*
- 9: /cgn2\_6/ptodata/1/pubpna/US09\_NEW\_PUB.seq.\*
- 10: /cgn2\_6/ptodata/1/pubpna/US09\_PUBCOMB.seq.\*
- 11: /cgn2\_6/ptodata/1/pubpna/US10\_NEW\_PUB.seq.\*
- 12: /cgn2\_6/ptodata/1/pubpna/US10\_PUBCOMB.seq.\*
- 13: /cgn2\_6/ptodata/1/pubpna/US60\_NEW\_PUB.seq.\*
- 14: /cgn2\_6/ptodata/1/pubpna/US60\_PUBCOMB.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	12	100.0	355	10	US-09-560-863-114
2	12	100.0	575	10	US-09-917-800A-992
3	12	100.0	1351	10	US-09-731-872-80
4	12	100.0	1474	10	US-09-731-872-139
5	12	100.0	1677	10	US-09-915-582-16
6	12	100.0	1802	10	US-09-799-777-104
7	12	100.0	1845	9	US-09-738-626-3487
8	12	100.0	1848	10	US-09-915-582-34
9	12	100.0	11009	10	US-09-845-583-1
10	11	91.7	118	10	US-09-867-701-6217
11	11	91.7	126	10	US-09-815-343-1511
12	11	91.7	200	9	US-09-900-714A-3
13	11	91.7	247	10	US-09-923-876-557
14	11	91.7	274	10	US-09-960-352-10276
15	11	91.7	286	9	US-09-841-157A-2
16	11	91.7	286	10	US-09-294-093B-2281
17	11	91.7	287	10	US-09-294-093B-3949
18	11	91.7	300	10	US-09-294-093B-3480
19	11	91.7	326	10	US-09-764-864-615

c 20	11	91.7	341	10	US-09-783-590-6885	Sequence 6885, Ap
c 21	11	91.7	350	10	US-09-815-343-1513	Sequence 1513, Ap
c 22	11	91.7	385	10	US-09-864-761-16659	Sequence 16659, A
c 23	11	91.7	405	9	US-09-738-626-2571	Sequence 2571, Ap
c 24	11	91.7	411	10	US-09-974-300-7207	Sequence 7207, Ap
c 25	11	91.7	415	10	US-09-834-975-417	Sequence 417, App
c 26	11	91.7	416	12	US-10-044-090-425	Sequence 425, App
c 27	11	91.7	434	10	US-09-960-352-1726	Sequence 1726, App
c 28	11	91.7	462	10	US-09-867-701-2628	Sequence 2628, Ap
c 29	11	91.7	474	9	US-10-025-380-1073	Sequence 1073, Ap
c 30	11	91.7	474	10	US-09-922-217-1073	Sequence 1073, Ap
c 31	11	91.7	474	10	US-09-833-263-1073	Sequence 1073, Ap
c 32	11	91.7	475	10	US-09-864-761-4882	Sequence 4882, Ap
c 33	11	91.7	481	10	US-09-998-598-25	Sequence 25, Appl
c 34	11	91.7	503	10	US-09-920-300A-1057	Sequence 1057, Ap
c 35	11	91.7	503	12	US-10-033-528-1057	Sequence 1057, Ap
c 36	11	91.7	551	10	US-09-962-832-3	Sequence 3, Appl
c 37	11	91.7	555	10	US-09-815-343-1539	Sequence 1539, Ap
c 38	11	91.7	555	10	US-09-998-598-1218	Sequence 1218, Ap
c 39	11	91.7	588	10	US-09-815-343-280	Sequence 280, App
c 40	11	91.7	609	10	US-09-815-242-6509	Sequence 6509, Ap
c 41	11	91.7	614	9	US-10-025-380-1075	Sequence 1075, Ap
c 42	11	91.7	614	10	US-09-922-217-1075	Sequence 1075, Ap
c 43	11	91.7	614	10	US-09-833-263-1075	Sequence 1075, Ap
c 44	11	91.7	666	9	US-09-738-626-706	Sequence 706, App
c 45	11	91.7	690	9	US-09-821-877-6	Sequence 6, Appl

ALIGNMENTS

RESULT 1  
US-09-560-863-114  
; Sequence 114, Application US/09560863  
; Patent No. US20020110809A1  
; GENERAL INFORMATION:  
; APPLICANT: Nehls, Michael C.  
; APPLICANT: Zambrowicz, Brian  
; APPLICANT: Sands, Arthur T.  
; TITLE OF INVENTION: No. US20020110809A1el Human Polynucleotides and the  
; FILE REFERENCE: Polypeptides Encoded Thereby  
; FILE REFERENCE: LEX-0018-USA  
; CURRENT APPLICATION NUMBER: US/09/560,863  
; PRIOR FILING DATE: 1999-04-28  
; PRIOR FILING DATE: 2000-04-30  
; NUMBER OF SEQ ID NOS: 1008  
; SOFTWARE: FASTSEQ for Windows Version 4.0  
; SEQ ID NO 114  
; LENGTH: 355  
; TYPE: DNA  
; ORGANISM: homo sapiens  
US-09-560-863-114

Query Match 100.0%; Score 12; DB 10; Length 355;  
Best Local Similarity 100.0%; Pred. No. 2.1e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGACGCA 12

Db 40 GTCGTGACGCA 51

RESULT 2

US-09-917-800A-992  
; Sequence 992, Application US/09917800A  
; Patent No. US20020119462A1  
; GENERAL INFORMATION:  
; APPLICANT: Mendrick, Donna  
; APPLICANT: Porter, Mark  
; APPLICANT: Johnson, Kory  
; APPLICANT: Castile, Arthur  
; APPLICANT: Elashoff, Michael

APPLICANT: Gene Logic, Inc.  
; TITLE OF INVENTION: Molecular Toxicology Modeling  
; FILE REFERENCE: 44921-5038-US  
; CURRENT APPLICATION NUMBER: US/09/917,800A  
; CURRENT FILING DATE: 2001-07-31  
; PRIOR APPLICATION NUMBER: US 60/222,040  
; PRIOR FILING DATE: 2000-07-31  
; PRIOR APPLICATION NUMBER: US 60/222,880  
; PRIOR FILING DATE: 2000-11-02  
; PRIOR APPLICATION NUMBER: US 60/290,029  
; PRIOR FILING DATE: 2001-05-11  
; PRIOR APPLICATION NUMBER: US 60/290,645  
; PRIOR FILING DATE: 2001-05-15  
; PRIOR APPLICATION NUMBER: US 60/292,336  
; PRIOR FILING DATE: 2001-05-22  
; PRIOR APPLICATION NUMBER: US 60/295,798  
; PRIOR FILING DATE: 2001-06-06  
; PRIOR APPLICATION NUMBER: US 60/297,457  
; PRIOR FILING DATE: 2001-06-13  
; PRIOR APPLICATION NUMBER: US 60/298,884  
; PRIOR FILING DATE: 2001-06-19  
; PRIOR APPLICATION NUMBER: US 60/303,459  
; PRIOR FILING DATE: 2001-07-09  
; NUMBER OF SEQ ID NOS: 1740  
; SOFTWARE: Patent In Ver. 2.1  
; SEQ ID NO 992  
; LENGTH: 575  
; TYPE: DNA  
; ORGANISM: Rattus norvegicus  
; FEATURE:  
; OTHER INFORMATION: Genbank Accession No. US20020119462A1 A1176942  
US-09-917-800A-992

Query Match 100.0%; Score 12; DB 10; Length 575;  
Best Local Similarity 100.0%; Pred. No. 2.le+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
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Db 68 GTCGTGCAGGCA 79

RESULT 3  
US-09-731-872-80  
; Sequence 80, Application US/09731872  
; Patent No. US20020102604A1  
; GENERAL INFORMATION:  
; APPLICANT: Dumas Milne Edwards, Jean Baptiste  
; APPLICANT: Bougueleret, Lydie  
; APPLICANT: Jobert, Severin  
; TITLE OF INVENTION: FULL-LENGTH HUMAN cDNAs ENCODING POTENTIALLY SECRETED PROTEINS  
; FILE REFERENCE: 78.US3.REG  
; CURRENT APPLICATION NUMBER: US/09/731,872  
; CURRENT FILING DATE: 2000-12-07  
; PRIOR APPLICATION NUMBER: US 60/169,629  
; PRIOR FILING DATE: 1999-12-08  
; PRIOR APPLICATION NUMBER: US 60/187,470  
; PRIOR FILING DATE: 2000-03-06  
; NUMBER OF SEQ ID NOS: 482  
; SOFTWARE: Patent.pm  
; SEQ ID NO 80  
; LENGTH: 1351  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 259..831  
; NAME/KEY: sig\_peptide  
; LOCATION: 259..375  
; OTHER INFORMATION: Von Heijne matrix  
; OTHER INFORMATION: score 5.809301698725  
; OTHER INFORMATION: seq FCVCVIAIGVVQA/LI  
US-09-731-872-80

Query Match 100.0%; Score 12; DB 10; Length 1351;  
Best Local Similarity 100.0%; Pred. No. 2.le+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
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Db 364 GTCGTGCAGGCA 375

RESULT 4  
US-09-731-872-139  
; Sequence 139, Application US/09731872  
; Patent No. US20020102604A1  
; GENERAL INFORMATION:  
; APPLICANT: Dumas Milne Edwards, Jean Baptiste  
; APPLICANT: Bougueleret, Lydie  
; APPLICANT: Jobert, Severin  
; TITLE OF INVENTION: FULL-LENGTH HUMAN cDNAs ENCODING POTENTIALLY SECRETED PROTEINS  
; FILE REFERENCE: 78.US3.REG  
; CURRENT APPLICATION NUMBER: US/09/731,872  
; CURRENT FILING DATE: 2000-12-07  
; PRIOR APPLICATION NUMBER: US 60/169,629  
; PRIOR FILING DATE: 1999-12-08  
; PRIOR APPLICATION NUMBER: US 60/187,470  
; PRIOR FILING DATE: 2000-03-06  
; NUMBER OF SEQ ID NOS: 482  
; SOFTWARE: Patent.pm  
; SEQ ID NO 139  
; LENGTH: 1474  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 16..471  
; NAME/KEY: sig\_peptide  
; LOCATION: 16..93  
; OTHER INFORMATION: Von Heijne matrix  
; OTHER INFORMATION: score 5.809301698725  
; OTHER INFORMATION: seq FCVCVIAIGVVQA/LI  
US-09-731-872-139

Query Match 100.0%; Score 12; DB 10; Length 1474;  
Best Local Similarity 100.0%; Pred. No. 2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
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Db 82 GTCGTGCAGGCA 93

RESULT 5  
US-09-915-582-16  
; Sequence 16, Application US/09915582  
; Patent No. US20020120103A1  
; GENERAL INFORMATION:  
; APPLICANT: Rosen et al.  
; TITLE OF INVENTION: 17 Human Secreted Proteins  
; FILE REFERENCE: PS723PI  
; CURRENT APPLICATION NUMBER: US/09/915,582  
; CURRENT FILING DATE: 2001-07-27  
; PRIOR APPLICATION NUMBER: PCT/US01/01431  
; PRIOR FILING DATE: 2001-01-17  
; PRIOR APPLICATION NUMBER: 60/179,065  
; PRIOR FILING DATE: 2000-01-31  
; PRIOR APPLICATION NUMBER: 60/180,628  
; PRIOR FILING DATE: 2000-02-04  
; PRIOR APPLICATION NUMBER: 60/231,968  
; PRIOR FILING DATE: 2000-09-12  
; NUMBER OF SEQ ID NOS: 97  
; SOFTWARE: Patent In Ver. 2.0  
; SEQ ID NO 16  
; LENGTH: 1677



; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-915-582-16

Query Match 100.0%; Score 12; DB 10; Length 1677;  
Best Local Similarity 100.0%; Pred. No. 2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
Db 200 GTCGTGCAGGCA 211  
|||||

## RESULT 6

US-09-799-777-104  
; Sequence 104, Application US/09799777  
; Patent No. US20020091244A1  
; GENERAL INFORMATION:

APPLICANT: Lal, Preeti  
Hillman, Jennifer L.  
Corley, Neil C.  
Guegler, Karl J.  
Baugh, Mariah  
Sather, Susan  
Shah, Purvi  
TITLE OF INVENTION: HUMAN SIGNAL PEPTIDE-CONTAINING PROTEINS  
NUMBER OF SEQUENCES: 154  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: INCYTE PHARMACEUTICALS, INC.  
STREET: 3174 PORTER DRIVE  
CITY: PALO ALTO  
STATE: CALIFORNIA  
COUNTRY: USA  
ZIP: 94304

## COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/799,777  
FILING DATE: 06-Mar-2001  
CLASSIFICATION: <Unknown>

## PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/09/002,485

## FILING DATE: &lt;Unknown&gt;

## ATTORNEY/AGENT INFORMATION:

NAME: BILLINGS, LUCY J.

REGISTRATION NUMBER: 36,749

REFERENCE/DOCKET NUMBER: PF-0459 US

## TELECOMMUNICATION INFORMATION:

TELEPHONE: (650) 855-0555

TELEFAX: (650) 845-4166

## INFORMATION FOR SEQ ID NO: 104:

## SEQUENCE CHARACTERISTICS:

LENGTH: 1802 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

## IMMEDIATE SOURCE:

LIBRARY: PROSTUT08

CLONE: 1653112

## SEQUENCE DESCRIPTION: SEQ ID NO: 104 :

US-09-799-777-104

Query Match 100.0%; Score 12; DB 10; Length 1802;  
Best Local Similarity 100.0%; Pred. No. 2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
Db 583 GTCGTGCAGGCA 594  
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## RESULT 7

US-09-738-626-3487  
; Sequence 3487, Application US/09738626  
; Publication No. US20020197605A1  
; GENERAL INFORMATION:

APPLICANT: NAKAGAWA, SATOSHI

APPLICANT: MIZOGUCHI, HIROSHI

APPLICANT: ANDO, SEIKO

APPLICANT: HAYASHI, MIKIRO

APPLICANT: OCHIAI, KEIKO

APPLICANT: YOKOI, HARUHIKO

APPLICANT: TATEISHI, NAKKO

APPLICANT: SENOH, AKIHIRO

APPLICANT: IKEDA, MASATO

APPLICANT: OZAKI, AKIO

TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES

FILE REFERENCE: 249-125

CURRENT APPLICATION NUMBER: US/09/738,626

CURRENT FILING DATE: 2000-12-18

PRIOR APPLICATION NUMBER: JP 99/377484

PRIOR FILING DATE: 1999-12-16

PRIOR APPLICATION NUMBER: JP 00/159162

PRIOR FILING DATE: 2000-04-07

PRIOR APPLICATION NUMBER: JP 00/280988

PRIOR FILING DATE: 2000-08-03

NUMBER OF SEQ ID NOS: 7059

SOFTWARE: PatentIn ver. 3.0

SEQ ID NO 3487

LENGTH: 1845

TYPE: DNA

ORGANISM: Corynebacterium glutamicum

US-09-738-626-3487

## Query Match

100.0%; Score 12; DB 9; Length 1845;

Best Local Similarity 100.0%; Pred. No. 2e+02;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12

Db 1139 GTCGTGCAGGCA 1150

## RESULT 8

US-09-915-582-34

; Sequence 34, Application US/09915582

; Patent No. US20020120103A1

; GENERAL INFORMATION:

APPLICANT: Rosen et al.

TITLE OF INVENTION: 17 Human Secreted Proteins

FILE REFERENCE: PS723p1

CURRENT APPLICATION NUMBER: US/09/915,582

CURRENT FILING DATE: 2001-07-27

PRIOR APPLICATION NUMBER: PCT/US01/01431

PRIOR FILING DATE: 2001-01-17

PRIOR APPLICATION NUMBER: 60/179,065

PRIOR FILING DATE: 2000-01-31

PRIOR APPLICATION NUMBER: 60/180,628

PRIOR FILING DATE: 2000-02-04

PRIOR APPLICATION NUMBER: 60/231,968

PRIOR FILING DATE: 2000-09-12

NUMBER OF SEQ ID NOS: 97

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 34

LENGTH: 1848

TYPE: DNA

ORGANISM: Homo sapiens

FEATURE:

NAME/KEY: SITE

LOCATION: (1796)

OTHER INFORMATION: n equals a,t,g, or c

US-09-915-582-34

Query Match 100.0%; Score 12; DB 10; Length 1848;  
Best Local Similarity 100.0%; Pred. No. 2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
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Db 378 GTCGTGCAGGCA 389

## RESULT 9

US-09-845-583-1/c  
; Sequence 1, Application US/09845583  
; Patent No. US20020142954A1  
; GENERAL INFORMATION:  
; APPLICANT: Burgeson, Robert  
; APPLICANT: Brunken, William Joseph  
; APPLICANT: Champlaud, Marie-France  
; APPLICANT: Hunter, Dale  
; TITLE OF INVENTION: LAMININ 15 AND USES THEREOF  
; FILE REFERENCE: 10287-056001  
; CURRENT APPLICATION NUMBER: US/09/845,583  
; CURRENT FILING DATE: 2001-04-30  
; PRIOR APPLICATION NUMBER: US 60/200,863  
; PRIOR FILING DATE: 2000-05-01  
; NUMBER OF SEQ ID NOS: 18  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 11009  
; TYPE: DNA  
; ORGANISM: Mus musculus  
US-09-845-583-1

Query Match 100.0%; Score 12; DB 10; Length 11009;  
Best Local Similarity 100.0%; Pred. No. 2e+02;  
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGCA 12  
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Db 9731 GTCGTGCAGGCA 9720

## RESULT 10

US-09-867-701-6217/c  
; Sequence 6217, Application US/09867701  
; Patent No. US20020132237A1  
; GENERAL INFORMATION:  
; APPLICANT: Aglate, Paul A.  
; APPLICANT: Jones, Robert  
; APPLICANT: Harlocker, Susan L.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY  
; TITLE OF INVENTION: AND DIAGNOSIS OF OVARIAN CANCER  
; FILE REFERENCE: 210121.497  
; CURRENT APPLICATION NUMBER: US/09/867,701  
; CURRENT FILING DATE: 2001-05-29  
; NUMBER OF SEQ ID NOS: 10912  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 6217  
; LENGTH: 118  
; TYPE: DNA  
; ORGANISM: Homo sapien  
US-09-867-701-6217

Query Match 91.7%; Score 11; DB 10; Length 118;  
Best Local Similarity 100.0%; Pred. No. 8.3e+02;  
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGC 11  
|||||  
Db 75 GTCGTGCAGGC 65

## RESULT 11

US-09-815-343-1511/c

; Sequence 1511, Application US/09815343  
; Patent No. US20010055596A1  
; GENERAL INFORMATION:  
; APPLICANT: Meagher, Madeleine  
; APPLICANT: Xu, Jiangchun E.  
; APPLICANT: King, Gordon E.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THERAPY AND  
; TITLE OF INVENTION: DIAGNOSIS OF COLON CANCER  
; FILE REFERENCE: 210121.504  
; CURRENT APPLICATION NUMBER: US/09/815,343  
; CURRENT FILING DATE: 2001-03-22  
; NUMBER OF SEQ ID NOS: 1556  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1511  
; LENGTH: 126  
; TYPE: DNA  
; ORGANISM: Homo sapien  
; FEATURE:  
; NAME/KEY: misc.feature  
; LOCATION: (1)...(126)  
; OTHER INFORMATION: n = A,T,C or G  
US-09-815-343-1511

Query Match 91.7%; Score 11; DB 10; Length 126;  
Best Local Similarity 100.0%; Pred. No. 8.3e+02;  
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCGTGCAGGC 11  
|||||  
Db 76 GTCGTGCAGGC 66

## RESULT 12

US-09-900-714A-3  
; Sequence 3, Application US/09900714A  
; Patent No. US20020162133A1  
; GENERAL INFORMATION:  
; APPLICANT: Allen, Keith D.  
; TITLE OF INVENTION: TRANSGENIC MICE CONTAINING MGLUR8  
; TITLE OF INVENTION: METABOTROPIC GLUTAMATE RECEPTOR GENE DISRUPTIONS  
; FILE REFERENCE: R-657  
; CURRENT APPLICATION NUMBER: US/09/900,714A  
; CURRENT FILING DATE: 2001-07-06  
; PRIOR APPLICATION NUMBER: US 60/216,252  
; PRIOR FILING DATE: 2000-07-06  
; PRIOR APPLICATION NUMBER: US 60/221,490  
; PRIOR FILING DATE: 2000-07-27  
; PRIOR APPLICATION NUMBER: US 60/262,138  
; PRIOR FILING DATE: 2001-01-16  
; PRIOR APPLICATION NUMBER: US 60/300,928  
; PRIOR FILING DATE: 2000-07-26  
; NUMBER OF SEQ ID NOS: 3  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 3  
; LENGTH: 200  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Targeting vector  
US-09-900-714A-3

Query Match 91.7%; Score 11; DB 9; Length 200;  
Best Local Similarity 100.0%; Pred. No. 8.3e+02;  
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCGTGCAGGCA 12  
|||||  
Db 61 TCGTGCAGGCA 71

## RESULT 13

US-09-923-876-557  
; Sequence 557, Application US/09923876

```

; Patent No. US20020013958A1
; GENERAL INFORMATION:
; APPLICANT: Lalgudi, Raghunath V.
; APPLICANT: Kamigaki, Laura Y. (Ito)
; APPLICANT: Sherman, Bradley K.
; TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES DERIVED FROM CORN SEEDLING
; FILE REFERENCE: PL-0012-1 CON
; CURRENT APPLICATION NUMBER: US/09/923,876
; CURRENT FILING DATE: 2001-08-06
; PRIOR APPLICATION NUMBER: 09/298,329
; PRIOR FILING DATE: 1999-04-21
; PRIOR APPLICATION NUMBER: 60/085,331
; PRIOR FILING DATE: 1998-05-05
; NUMBER OF SEQ ID NOS: 6332
; SOFTWARE: PERL Program
; SEQ ID NO 557
; LENGTH: 247
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: misc-feature
; OTHER INFORMATION: Incyte ID No. US20020013958A1 700157144H1
US-09-923-876-557

```

```

Query Match          91.7%; Score 11; DB 10; Length 247;
Best Local Similarity 100.0%; Pred. No. 8.3e+02;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 2 TCGTGCAGGCA 12
    |||||
Db 170 TCGTGCAGGCA 180

```

# RESULT 14

```

US-09-960-352-10276/c
; Sequence 10276, Application US/09960352
; Patent No. US20020137139A1
; GENERAL INFORMATION:
; APPLICANT: Warren, Wesley C.
; APPLICANT: Tao, Nengbing
; APPLICANT: Byatt, John C.
; APPLICANT: Mathialagan, Nagappan
; TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH LACTATION AND
; FILE REFERENCE: 16511.006/37-21(10298)C
; CURRENT APPLICATION NUMBER: US/09/960,352
; CURRENT FILING DATE: 2001-09-24
; NUMBER OF SEQ ID NOS: 15112
; SEQ ID NO 10276
; LENGTH: 274
; TYPE: DNA
; ORGANISM: Bos taurus
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (19)
; OTHER INFORMATION: unsure at all n locations
; OTHER INFORMATION: Clone ID: 44-LIB3058-056-Q1-K1-C8
US-09-960-352-10276

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```

Query Match          91.7%; Score 11; DB 10; Length 274;
Best Local Similarity 100.0%; Pred. No. 8.2e+02;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 GTCGTGCAGGC 11
    |||||
Db 57 GTCGTGCAGGC 47

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# RESULT 15

```

US-09-841-157A-2
; Sequence 2, Application US/09841157A
; Publication No. US20020192648A1
; GENERAL INFORMATION:

```

```

; APPLICANT: NISHIGAKI, KOICHI
; APPLICANT: TAKASAWA, TSUTOMU
; APPLICANT: HAMANO, KEIICHI
; TITLE OF INVENTION: METHODS OF IDENTIFYING AN ORGANISM BASED ON ITS GENOTYPE
; FILE REFERENCE: 12637/P66602USO
; CURRENT APPLICATION NUMBER: US/09/841,157A
; CURRENT FILING DATE: 2001-04-25
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 286
; TYPE: DNA
; ORGANISM: Unknown Organism
; FEATURE:
; OTHER INFORMATION: Description of Unknown Organism: Illustrative standard
US-09-841-157A-2

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Query Match          91.7%; Score 11; DB 9; Length 286;
Best Local Similarity 100.0%; Pred. No. 8.2e+02;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 2 TCGTGCAGGCA 12
    |||||
Db 180 TCGTGCAGGCA 190

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Search completed: January 4, 2003, 01:06:18
Job time : 6.05047 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:46 ; Search time 216.151 Seconds  
(without alignments)  
3231.380 Million cell updates/sec

Title: US-09-787-562-11

Perfect score: 24

Sequence: 1 tctagtgtctgtcagcatcatgt 24

Scoring table: IDENTITY\_NUC

Gapop 10.0, Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: gb\_ba.\*

2: gb\_htg.\*

3: gb\_in.\*

4: gb\_om.\*

5: gb\_ov.\*

6: gb\_pat.\*

7: gb\_ph.\*

8: gb\_pl.\*

9: gb\_pr.\*

10: gb\_ro.\*

11: gb\_sts.\*

12: gb\_sy.\*

13: gb\_un.\*

14: gb\_vi.\*

15: em\_ba.\*

16: em\_fun.\*

17: em\_in.\*

18: em\_mu.\*

19: em\_mu.\*

20: em\_om.\*

21: em\_or.\*

22: em\_ov.\*

23: em\_pat.\*

24: em\_ph.\*

25: em\_pl.\*

26: em\_ro.\*

27: em\_sts.\*

28: em\_un.\*

29: em\_vi.\*

30: em\_htg\_hum.\*

31: em\_htg\_inv.\*

32: em\_htg\_other.\*

33: em\_htg\_mus.\*

34: em\_htg\_pin.\*

35: em\_htg\_rod.\*

36: em\_htg\_mam.\*

37: em\_htg\_vrt.\*

38: em\_sy.\*

39: em\_htgo\_hum.\*

40: em\_htgo\_mus.\*

41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	24	100.0	24	6	AX023669 Sequence
2	24	100.0	237	6	AX023667 Sequence
3	18.8	78.3	183342	2	AC129440 Rattus no
4	18.2	75.8	64672	2	AC126356 Homo sapi
5	18.2	75.8	103432	2	AC094981 Rattus no
6	18.2	75.8	182943	9	AC016251 Homo sapi
7	18.2	75.8	200078	10	AL591064 Mouse DNA
8	18.2	75.8	229662	2	AC125109 Mus muscu
9	17.8	74.2	1768	9	HSAMPD3S24
10	17.8	74.2	40520	1	AC090969 Staphyloc
11	17.8	74.2	69320	2	AC113035 Mus muscu
12	17.8	74.2	152631	2	AC048373 Homo sapi
13	17.8	74.2	177429	9	AC021914 Homo sapi
14	17.8	74.2	177488	9	AP0002378 Homo sapi
15	17.8	74.2	204140	9	AP000760 Homo sapi
16	17.8	74.2	272850	1	AP0010976 Homo sapi
17	17.8	74.2	272850	1	AP004828 Staphyloc
18	17.8	74.2	291150	1	AP003135 Staphyloc
19	17.8	74.2	342600	1	AP003363 Staphyloc
20	17.6	73.3	6237	9	D86978 Human mRNA
21	17.6	73.3	82642	2	AC111397 Rattus no
22	17.6	73.3	110000	2	AC095071_3 Continuation (4 of
23	17.6	73.3	168089	2	AC113849 Rattus no
24	17.6	73.3	183977	2	AC097956 Rattus no
25	17.6	73.3	197837	9	AC093107 Homo sapi
26	17.6	73.3	207585	2	AC073779 Mus muscu
27	17.6	73.3	235218	2	AC074208 Mus muscu
28	17.6	73.3	247593	2	AC103319 Rattus no
29	17.4	72.5	72	6	AX023690 Sequence
30	17.4	72.5	72	6	AX023692 Sequence
31	17.2	71.7	1676	1	ECONEUC
32	17.2	71.7	59634	9	AC062028 Homo sapi
33	17.2	71.7	61796	2	AC020875 Mus muscu
34	17.2	71.7	95560	2	AC127620 Rattus no
35	17.2	71.7	102345	9	AL445230 Human DNA
36	17.2	71.7	120739	2	AL807782 Homo sapi
37	17.2	71.7	170677	2	AL138932 Homo sapi
38	17.2	71.7	170965	2	AC108288 Rattus no
39	17.2	71.7	175758	9	AC098825 Homo sapi
40	17.2	71.7	175957	2	AL691501 Mus muscu
41	17.2	71.7	180418	9	AC093875 Homo sapi
42	17.2	71.7	183338	2	AC120970 Rattus no
43	17.2	71.7	185425	2	AC124837 Rattus no
44	17.2	71.7	185532	10	AC121883 Mus muscu
45	17.2	71.7	196203	9	AL355355 Human DNA

# ALIGNMENTS

RESULT 1  
AX023669

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

AX023669 Sequence 11 from Patent WO0017371. 24 bp DNA Linear PAT 16-SEP-2000

AX023669

AX023669.1 GI:10184030

synthetic construct.

synthetic construct

artificial sequences.

1 (bases 1 to 24)

Binley,K.M. and Naylor,S.

Polynucleotide constructs and uses thereof

Patent: WO 0017371-A 11 30-MAR-2000;

BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD

FEATURES		(GB)	
source		Location/Qualifiers	
		1. .24	
		/organism="synthetic construct"	
		/db xref="taxon:32630"	
		/note="Spacer"	
BASE COUNT		4 a 5 c 7 g 8 t	
ORIGIN			
Query Match		100.0%; Score 24; DB 6; Length 24;	
Best Local Similarity		100.0%; Pred. No. 0.045;	
Matches		24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
QY		1 TCTAGTGTGTCGAGGCATCTAGT 24	
Db		1 TCTAGTGTGTCGAGGCATCTAGT 24	
RESULT 2			
AX023667		237 bp DNA linear PAT 15-SEP-2000	
LOCUS		Sequence 9 from Patent WO0017371.	
DEFINITION		AX023667	
ACCESSION		AX023667	
VERSION		AX023667.1 GI:10184028	
KEYWORDS		synthetic construct.	
SOURCE		artificial construct.	
ORGANISM		1 (bases 1 to 237)	
REFERENCE		Binley K.M. and Naylor S.	
AUTHORS		Polynucleotide constructs and uses thereof	
TITLE		Patent: WO 0017371-A 9 30-MAR-2000;	
JOURNAL		BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD	
FEATURES		(GB)	
source		Location/Qualifiers	
		1. .237	
		/organism="synthetic construct"	
		/db xref="taxon:32630"	
		/note="OBHrel1"	
BASE COUNT		43 a 82 c 56 g 56 t	
ORIGIN			
Query Match		100.0%; Score 24; DB 6; Length 237;	
Best Local Similarity		100.0%; Pred. No. 0.041;	
Matches		24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
QY		1 TCTAGTGTGTCGAGGCATCTAGT 24	
Db		25 TCTAGTGTGTCGAGGCATCTAGT 48	
RESULT 3			
AC129440		183342 bp DNA linear HTG 30-JUL-2002	
LOCUS		Rattus norvegicus clone CH230-112K14, *** SEQUENCING IN PROGRESS	
DEFINITION		***, 65 unordered pieces.	
ACCESSION		AC129440	
VERSION		AC129440.1 GI:22004125	
KEYWORDS		HTG; HTGS_PHASE1.	
SOURCE		Norway rat.	
ORGANISM		Rattus norvegicus	
REFERENCE		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
AUTHORS		Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae;	
		Rattus.	
		1 (bases 1 to 183342)	
		Muzny D.M., Adams C., Adio-Oduola B., Ali-oshan F.R., Allen C.,	
		Alsbrooks S.L., Anaratunge H.C., Are J.R., Ayele M., Banks T.,	
		Barbaria J., Benton J., Binage K., Blankenburg K., Bonnin D.,	
		Bouck J., Bowie S., Brieva M., Brown E., Brown M., Bryant N.P.,	
		Buhay C., Burch P., Burkett C., Burrell K.L., Byrd N.C.,	
		Carroll T.F., Carter M., Cavazos S.R., Chacko J., Chavez D.,	
		Chen G., Chen K., Chen Z., Chowdhry I., Christopoulos C.,	
		Cleveland C.D., Cox C., Coyle M.D., Dathorne S.K., David R.,	

Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A., Delaney, K.R., Delgado, O., Denn, A.L., Ding, Y., Dinh, H.H., Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J., Earnhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Escotto, M., Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R., Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K., Harris, C., Harris, K., Hart, M., Havlak, P., Hawes, A., Hernandez, J., Hernandez, O., Hodgson, A., Hogue, M., Holloway, C., Hollins, B., Homi, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Jackson, L.E., Jacobson, B., Jia, J., Johnson, R., Jolivet, S., Joudah, S., Karlsson, E., Kelly, S., Khan, U., King, L., Korvan, J., Kovar, C., Kratoch, J., Kureshi, A., Landry, N., Leal, B., Lewis, L.C., Lewis, L., Li, J., Li, Z., Lichtarge, O., Lieu, C., Liu, J., Liu, W., Loulseged, H., Lozano, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J., Maheshwari, M., Mapua, P., Martin, R., Martindale, A., Martinez, E., Massey, B., Mawhney, E., McLeod, M.P., Meador, M., Mei, G., Mettler, M., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Morgan, M., Morris, S., Moser, M., Neal, D., Newton, N., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokwos, S., Ogih, M., Okwuonu, G., Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Picken, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y., Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savery, G., Scherer, S., Scott, G., Shen, H., Shoostari, N., Sisson, I., Sodergren, E., Sonaik, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Tabor, P., Tamerisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telifrod, B., Thomas, N., Thomas, S., Usmani, K., Vasquez, L., Vera, V., Villalón, D., Vinson, R., Wang, Q., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleczyk, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D., Weinstein, G., and Gibbs, R.

Direct Submission  
Unpublished  
2 (bases 1 to 183342)  
Worley, K.C.  
Direct Submission  
Submitted (30-JUL-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA  
----- Genome Center  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)  
----- Project Information  
Center project name: GKTY  
Center clone name: CH230-112K14  
----- Summary Statistics  
Sequencing vector: Plasmid;  
Chemistry: Dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.990329  
Consensus quality: 130157 bases at least Q40  
Consensus quality: 135959 bases at least Q30  
Consensus quality: 140433 bases at least Q20  
-----  
\* NOTE: Estimated insert size may differ from sequence length (see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently consists of 65 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.  
\* 1 1012: contig of 1012 bp in length  
\* 1013 1112: gap of unknown length  
\* 1113 2140: contig of 1028 bp in length  
\* 2141 2240: gap of unknown length  
\* 2241 3744: contig of 1504 bp in length  
\* 3745 3844: gap of unknown length  
\* 3845 4961: contig of 1117 bp in length

```

* 4962 5061: gap of unknown length
* 5062 6356: contig of 1295 bp in length
* 6456: gap of unknown length
* 6457 8020: contig of 1564 bp in length
* 8021 8120: gap of unknown length
* 8121 9729: contig of 1609 bp in length
* 9730 9829: gap of unknown length
* 9830 10968: contig of 1139 bp in length
* 10969 11068: gap of unknown length
* 11070 12102: contig of 1034 bp in length
* 12103 12202: gap of unknown length
* 12204 13464: contig of 1262 bp in length
* 13465 13564: gap of unknown length
* 13565 15019: contig of 1455 bp in length
* 15020 15119: gap of unknown length
* 15120 16277: contig of 1158 bp in length
* 16278 16377: gap of unknown length
* 16379 17598: contig of 1221 bp in length
* 17599 17698: gap of unknown length
* 17699 19126: contig of 1428 bp in length
* 19127 19226: gap of unknown length
* 19228 20838: contig of 1612 bp in length
* 20839 22282: contig of 1344 bp in length
* 22283 23806: contig of 1424 bp in length
* 23807 23906: gap of unknown length
* 23907 25005: contig of 1099 bp in length
* 25006 25105: gap of unknown length
* 25106 26897: contig of 1792 bp in length
* 26898 26997: gap of unknown length
* 26998 29011: contig of 2014 bp in length
* 29012 29111: gap of unknown length
* 29112 30662: contig of 1551 bp in length
* 30663 30762: gap of unknown length
* 30763 32775: contig of 2013 bp in length
* 32776 32875: gap of unknown length
* 32876 34108: contig of 1233 bp in length
* 34109 34208: gap of unknown length
* 34209 35893: contig of 1685 bp in length
* 35894 35993: gap of unknown length
* 35994 38034: contig of 2041 bp in length
* 38035 38134: gap of unknown length
* 38135 39356: contig of 1222 bp in length
* 39357 39456: gap of unknown length
* 39457 41734: contig of 2278 bp in length
* 41735 41834: gap of unknown length
* 41835 43106: contig of 1272 bp in length
* 43107 43206: gap of unknown length
* 43207 44719: contig of 1513 bp in length
* 44720 46402: contig of 1583 bp in length
* 46403 46502: gap of unknown length
* 46503 48677: contig of 2175 bp in length
* 48678 48777: gap of unknown length
* 48779 50549: contig of 1772 bp in length
* 50550 50649: gap of unknown length
* 50650 51975: contig of 1326 bp in length
* 51976 52075: gap of unknown length
* 52076 54622: contig of 2547 bp in length
* 54623 54722: gap of unknown length
* 54723 57428: contig of 2706 bp in length
* 57429 60703: contig of 3175 bp in length
* 60704 60803: gap of unknown length
* 60804 62969: contig of 2166 bp in length
* 62970 63069: gap of unknown length
* 63070 65319: contig of 2250 bp in length
* 65320 65419: gap of unknown length
* 65420 68626: contig of 3207 bp in length
* 68627 71376: gap of unknown length
* 71377 71476: gap of unknown length

* 71477 73563: contig of 2087 bp in length
* 73564 73663: gap of unknown length
* 73664 76959: contig of 3295 bp in length
* 76960 77059: gap of unknown length
* 77060 79870: contig of 2812 bp in length
* 79871 79970: gap of unknown length
* 79971 83050: contig of 3080 bp in length
* 83051 83150: gap of unknown length
* 83151 86495: contig of 3345 bp in length
* 86496 86596: gap of unknown length
* 86597 89682: contig of 3086 bp in length
* 89683 92489: contig of 2708 bp in length
* 92490 92589: gap of unknown length
* 92590 95401: contig of 2812 bp in length
* 95402 95501: gap of unknown length
* 95502 98592: contig of 3091 bp in length
* 98593 101719: gap of unknown length
* 101720 101819: contig of 3027 bp in length
* 101820 104542: contig of 2723 bp in length
* 104543 104642: gap of unknown length
* 104643 108398: contig of 3756 bp in length
* 108399 108498: gap of unknown length
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Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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DEFINITION AC126356
ACCESSION  AC126356
VERSION    AC126356.2 GI:22004353
KEYWORDS   HTG; HTGS_PHASE0.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 64672)
AUTHORS   Birren,B., Nussbaum,C. and Lander,E.
TITLE     Homo sapiens chromosome 17, clone CTD-2541111
JOURNAL   Unpublished
REFERENCE  2 (bases 1 to 64672)
AUTHORS   Birren,B., Nussbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
            Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B.,
            Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
            Cook,A., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
            Fato,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
            Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
            Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
            Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
            Liu,G., MacLean,C., Macdonald,P., Major,J., Matthews,C.,
            McCarthy,M., Meldrum,J., Meneus,L., Milhova,T., Mlenga,V.,
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            O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,

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TITLE  
JOURNALREFERENCE  
AUTHORS

Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission  
Submitted (05-JUL-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
3 (bases 1 to 64672)

Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhgalter, B., Canarata, J., Chang, J., Chazaro, P., Choepel, Y., Collymore, A., Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission  
Submitted (30-JUL-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Jul 30, 2002 this sequence version replaced gi:21699304.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997)  
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

TITLE  
JOURNAL

## COMMENT

Center: Whitehead Institute/ MIT Center for Genome Research  
Center code: WIBR  
Web site: <http://www-seq.wi.mit.edu>  
Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)  
----- Project Information  
Center project name: L27688  
Center clone name: 2541\_I\_11  
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\* NOTE: This record contains 78 individual  
\* sequencing reads that have not been assembled into  
\* contigs. Runs of N are used to separate the reads  
\* and the order in which they appear is completely  
\* arbitrary. Low-pass sequence sampling is useful for  
\* identifying clones that may be gene-rich and allows  
\* overlap relationships among clones to be deduced.  
\* However, it should not be assumed that this clone  
\* will be sequenced to completion. In the event that  
\* the record is updated, the accession number will  
\* be preserved.

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\* 705 804: gap of 100 bp  
\* 805 1540: contig of 736 bp in length  
\* 1541 1640: gap of 100 bp  
\* 1641 2370: contig of 730 bp in length  
\* 2371 2470: gap of 100 bp  
\* 2471 3203: contig of 733 bp in length  
\* 3204 3303: gap of 100 bp  
\* 3304 4019: contig of 716 bp in length  
\* 4020 4119: gap of 100 bp  
\* 4120 4870: contig of 751 bp in length  
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\* 4971 5690: contig of 720 bp in length  
\* 5691 5790: gap of 100 bp  
\* 5791 6514: contig of 724 bp in length  
\* 6515 6614: gap of 100 bp  
\* 6615 7363: contig of 749 bp in length  
\* 7364 7463: gap of 100 bp

7464 8203: contig of 740 bp in length  
\* 8204 8303: gap of 100 bp  
\* 8304 9028: contig of 725 bp in length  
\* 9029 9128: gap of 100 bp  
\* 9129 9837: contig of 729 bp in length  
\* 9838 9957: gap of 100 bp  
\* 9958 10683: contig of 726 bp in length  
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\* 29106 29842: contig of 737 bp in length  
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\* 32337 32436: gap of 100 bp  
\* 32437 33163: contig of 727 bp in length  
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\* 33995 34094: gap of 100 bp  
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\* 35759 36505: contig of 747 bp in length  
\* 36506 36605: gap of 100 bp  
\* 36606 37335: contig of 730 bp in length  
\* 37336 37435: gap of 100 bp  
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10142 11400: contig of 1259 bp in length  
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Best Local Similarity 87.0%; Pred. No. 57;

Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CTAGTGTCTGTCAGGCATCTAGT 24

Db 64816 CGAGTGTCTGTCAGGCATATAGT 64838

# RESULT 6

AC016251

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

AC016251 182943 bp DNA linear PRI 05-FEB-2002  
 Homo sapiens chromosome 15, clone RP11-759A24, complete sequence.

AC016251 GI:18139499

HTG.

Homo sapiens

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 182943)

Birren,B., Linton,L., Nusbaum,C. and Lander,E.

Homo sapiens chromosome 15, clone RP11-759A24

Unpublished

2 (bases 1 to 182943)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,  
 Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhvalter,B.,  
 Brown,A., Castle,A., Colangelo,M., Collins,S., Collamore,A.,  
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 Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,  
 Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

Direct Submission

Submitted (23-NOV-1999) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

3 (bases 1 to 182943)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,  
 Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Boukhvalter,B.,  
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Landers, T., Lehoczy, J., Levine, R., Liu, G., MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrum, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schnupack, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission  
Submitted (05-FEB-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Jan 12, 2002 this sequence version replaced gi:17646997.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997)  
<http://ftp.genome.washington.edu/RN/RepeatMasker.html>

----- Genome Center  
Center: Whitehead Institute/ MIT Center for Genome Research  
Center code: WIBR  
Web site: <http://www-seq.wi.mit.edu>  
Contact: [sequence.submissions@genome.wi.mit.edu](mailto:sequence.submissions@genome.wi.mit.edu)  
----- Project Information  
Center project name: L4985  
Center clone name: 759\_A\_24  
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Query Match
Best Local Similarity 75.8%; Score 18.2; DB 9; Length 182943;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CTAGTGTCTGTCAGGCATCTAGT 24
||||| ||||| ||||| ||||| |||||
Db 84216 CTACTGTAGTGCAGGCATCTAAT 84238

RESULT 7
AL591064
LOCUS
DEFINITION
AL591064 Mouse DNA sequence from clone RP23-395E18 on chromosome 2, complete
sequence.
ACCESSION
AL591064
VERSION
AL591064.11 GI:20068460
KEYWORDS
HTG.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1
Kay, M.
Submitted (04-APR-2002) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquerry@sanger.ac.uk
On Apr 7, 2002 this sequence version replaced gi:17644288.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em: EMBL; Sw:
SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP
database can be found at
http://www.sanger.ac.uk/projects/C_elegans/wormpep
from the RPI-23 Mouse PAC Library
constructed by the group of Pieter de Jong.
For further details see http://www.chori.org/bacpac/home.htm
VECTOR: pBAC3.6.
Location/Qualifiers
1..200078
/organism="Mus musculus"
/db_xref="taxon:10090"
/chromosome="2"
/clone="RP23-395E18"
/clone_lib="RPI-23"
50890 a 49240 c 50588 g 49360 t

BASE COUNT
ORIGIN

```

```

Query Match
Best Local Similarity 75.8%; Score 18.2; DB 10; Length 200078;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CTAGTGTCTGTCAGGCATCTAGT 24
||||| ||||| ||||| ||||| |||||
Db 9792 CTAGTGTAGTGTGCATCTAGT 9814

RESULT 8
AC125109
LOCUS
DEFINITION
AC125109 Mus musculus chromosome UNK clone RP24-232H5, WORKING DRAFT
SEQUENCE, 5 unordered pieces.
ACCESSION
AC125109
VERSION
AC125109.1 GI:21490585
KEYWORDS
HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1
McPherson, J.D. and Waterston, R.H.
The sequence of Mus musculus clone
Unpublished
REFERENCE
2 (bases 1 to 229662)
McPherson, J.D. and Waterston, R.H.
Direct Submission
Submitted (20-JUN-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA

COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
Contact: submissions@wustl.wustl.edu
----- Project Information -----
Center project name: M_B80232H05
----- Summary Statistics -----
Sequencing vector: M13; 0%
Chemistry: Dye-primer ET; 0% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 227185 bases at least Q40
Consensus quality: 227749 bases at least Q30
Consensus quality: 228011 bases at least Q20
Insert size: 163000; agarose-fp
Quality coverage: 17.07 in Q20 bases; agarose-fp
Quality coverage: 12.64 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 5 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 19665: contig of 19665 bp in length
* 19666 19765: gap of unknown length
* 19766 85228: contig of 65463 bp in length
* 85229 85328: gap of unknown length
* 85329 125046: contig of 39718 bp in length
* 125047 125146: gap of unknown length
* 125147 171694: contig of 46548 bp in length
* 171695 171795: gap of unknown length
* 171795 229662: contig of 57868 bp in length.
* Location/Qualifiers
1..229662

```

FEATURES  
source

/organism="Mus musculus"		/number=9	
/db_xref="taxon:10090"		1223. >1768	
/chromosome="UNK"		/gene="AMPD3"	
/clone="RP24-232H5"		/number=9	
1. .19665		379 a 459 c 496 g 434 t	
/note="assembly_name:Contig10"			
19766. .85228			
/note="assembly_name:Contig11"			
85329. .125046			
/note="assembly_name:Contig12"			
125147. .171694			
/note="assembly_name:Contig13"			
171795. .229662			
/note="assembly_name:Contig14"			
73610 a 42903 c 42981 g 69761 t		407 others	
BASE COUNT			
ORIGIN			
Query Match		75.8%; Score 18.2; DB 2; Length 229662;	
Best Local Similarity		87.0%; Pred. No. 55;	
Matches		20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;	
QY		1 TCTAGTCTGTCGAGCATCTAG 23	
Db		201681 TTATGTCAGTCGAGCATCTAG 201703	
RESULT 9			
HSAMPD3S24/c			
LOCUS		1768 bp DNA linear PRI 11-JUL-1996	
DEFINITION		Human AMP deaminase (AMPD3) gene, exon 8 and 9.	
ACCESSION		U29917	
VERSION		U29917.1 GI:1002652	
KEYWORDS			
SEGMENT		24 of 32	
SOURCE		Homo sapiens.	
ORGANISM		Homo sapiens.	
REFERENCE		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
AUTHORS		Mahnke-Zizelman,D.K., Eddy,R., Shows,T.B. and Sabina,R.L.	
TITLE		Characterization of the human AMPD3 gene reveals that 5' exon usage is subject to transcriptional control by three tandem promoters and alternative splicing	
JOURNAL		Biochim. Biophys. Acta 1306 (1), 75-92 (1996)	
MEDLINE		96201708	
PUBMED		8611627	
REFERENCE		2 (bases 1 to 1768)	
AUTHORS		Sabina,R.L.	
TITLE		Direct Submission	
JOURNAL		Submitted (22-JUN-1995) Richard L. Sabina, Biochemistry, Medical College of Wisconsin, 8701 Watertown Plank Road, Milwaukee, WI 53226, USA	
COMMENT		approximately 100 bp to next reported sequence, GenBank Accession Number U29918	
FEATURES		Location/Qualifiers	
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		/chromosome="11"	
		/map="11p13-pter"	
		/cell_type="T-lymphocyte, cytotoxic"	
		/clone_lib="RPMI 8402, lambda 2001 library of R. Baer"	
		<1. .389	
		/gene="AMPD3"	
		/number=7	
		390. .521	
		/gene="AMPD3"	
		/number=8	
		522. .1058	
		/gene="AMPD3"	
		/number=8	
		1059. .1222	
		/gene="AMPD3"	
intron			
exon			
intron			
exon			

LOCUS AC113035 69320 bp DNA linear HTG 13-MAY-2002  
 DEFINITION Mus musculus clone RP23-222J21, LOW-PASS SEQUENCE SAMPLING.  
 ACCESSION AC113035  
 VERSION AC113035.2 GI:20531939  
 KEYWORDS HTG; HTGS\_PHASE0.  
 SOURCE Mus musculus.  
 ORGANISM Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 1 (bases 1 to 69320)  
 Birren, B., Linton, L., Nusbaum, C. and Lander, E.  
 Unpublished  
 2 (bases 1 to 69320)  
 Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,  
 Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Boukhgalter, B.,  
 Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B.,  
 Choepel, Y., Colangelo, M., Collins, S., Collamore, A., Cook, A.,  
 Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S.,  
 Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S.,  
 Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,  
 Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,  
 Kanat, A., Karatas, A., Kells, C., LaRocque, K., Lamazares, R.,  
 Landers, T., Lehoczy, J., Levine, R., Liu, G., MacLean, C.,  
 Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M.,  
 McEwan, P., McKernan, K., Meldrim, J., Meneus, L., Mihova, T.,  
 Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,  
 Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,  
 Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C.,  
 Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J.,  
 Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S.,  
 Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,  
 Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,  
 Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H.,  
 Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,  
 Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.  
 Direct Submission  
 Submitted (25-FEB-2002) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA  
 3 (bases 1 to 69320)  
 Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,  
 Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L.,  
 Boukhgalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J.,  
 Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collamore, A.,  
 Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S.,  
 Faro, S., Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S.,  
 Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,  
 Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,  
 Kanat, A., Karatas, A., Kells, C., LaRocque, K., Lamazares, R.,  
 Landers, T., Lehoczy, J., Levine, R., Lindblad-Toh, K., Liu, G.,  
 MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C.,  
 McCarthy, M., McEwan, P., McKernan, K., Meldrim, J., Meneus, L.,  
 Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R.,  
 Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,  
 Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V.,  
 Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,  
 Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R.,  
 Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,  
 Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,  
 Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H.,  
 Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,  
 Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.  
 Direct Submission  
 Submitted (13-MAY-2002) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA  
 On May 13, 2002 this sequence version replaced gi:18875122.  
 All repeats were identified using RepeatMasker:  
 Smit, A.F.A. & Green, P. (1996-1997)  
 http://ftp.genome.washington.edu/RM/RepeatMasker.html  
 ----- Genome Center  
 Center: Whitehead Institute/ MIT Center for Genome Research  
 Center code: WIBR  
 Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu  
 ----- Project Information  
 Center project name: L23562  
 Center clone name: 222\_J\_21  
 -----  
 \* NOTE: This record contains 87 individual  
 \* sequencing reads that have not been assembled into  
 \* contigs. Runs of N are used to separate the reads  
 \* and the order in which they appear is completely  
 \* arbitrary. Low-pass sequence sampling is useful for  
 \* identifying clones that may be gene-rich and allows  
 \* overlap relationships among clones to be deduced.  
 \* However, it should not be assumed that this clone  
 \* will be sequenced to completion. In the event that  
 \* the record is updated, the accession number will  
 \* be preserved.  
 \* 1 703: contig of 703 bp in length  
 \* 704 803: gap of 100 bp  
 \* 804 1508: contig of 705 bp in length  
 \* 1509 1608: gap of 100 bp  
 \* 1609 2275: contig of 667 bp in length  
 \* 2276 2375: gap of 100 bp  
 \* 2376 3084: contig of 709 bp in length  
 \* 3085 3184: gap of 100 bp  
 \* 3185 3892: contig of 708 bp in length  
 \* 3893 3992: gap of 100 bp  
 \* 3993 4678: contig of 686 bp in length  
 \* 4679 4778: gap of 100 bp  
 \* 4779 5470: contig of 692 bp in length  
 \* 5471 5570: gap of 100 bp  
 \* 5571 6259: contig of 689 bp in length  
 \* 6260 6359: gap of 100 bp  
 \* 6360 7049: contig of 690 bp in length  
 \* 7050 7149: gap of 100 bp  
 \* 7150 7834: contig of 685 bp in length  
 \* 7835 7934: gap of 100 bp  
 \* 7935 8650: contig of 716 bp in length  
 \* 8651 8750: gap of 100 bp  
 \* 8751 9448: contig of 698 bp in length  
 \* 9449 9548: gap of 100 bp  
 \* 9549 10262: contig of 714 bp in length  
 \* 10263 10362: gap of 100 bp  
 \* 10363 11069: contig of 707 bp in length  
 \* 11070 11169: gap of 100 bp  
 \* 11170 11807: contig of 638 bp in length  
 \* 11808 11907: gap of 100 bp  
 \* 11908 12576: contig of 669 bp in length  
 \* 12577 12676: gap of 100 bp  
 \* 12677 13371: contig of 695 bp in length  
 \* 13372 13471: gap of 100 bp  
 \* 13472 14184: contig of 713 bp in length  
 \* 14185 14284: gap of 100 bp  
 \* 14285 14990: contig of 706 bp in length  
 \* 14991 15090: gap of 100 bp  
 \* 15091 15791: contig of 701 bp in length  
 \* 15792 15891: gap of 100 bp  
 \* 15892 16604: contig of 713 bp in length  
 \* 16605 16704: gap of 100 bp  
 \* 16705 17407: contig of 703 bp in length  
 \* 17408 17507: gap of 100 bp  
 \* 17508 18207: contig of 700 bp in length  
 \* 18208 18307: gap of 100 bp  
 \* 18308 19012: contig of 705 bp in length  
 \* 19013 19112: gap of 100 bp  
 \* 19113 19799: contig of 687 bp in length  
 \* 19800 19899: gap of 100 bp  
 \* 19900 20594: contig of 695 bp in length  
 \* 20595 20694: gap of 100 bp  
 \* 20695 21386: contig of 692 bp in length  
 \* 21387 21486: gap of 100 bp  
 \* 21487 22178: contig of 692 bp in length  
 \* 22179 22278: gap of 100 bp  
 \* 22279 22983: contig of 705 bp in length

```

* 22984 23083: gap of 100 bp
* 23084 23700: contig of 617 bp in length
* 23701 23800: gap of 100 bp
* 23801 24512: contig of 712 bp in length
* 24513 24612: gap of 100 bp
* 24613 25318: contig of 706 bp in length
* 25319 25418: gap of 100 bp
* 25419 26133: contig of 715 bp in length
* 26134 26233: gap of 100 bp
* 26234 26953: contig of 720 bp in length
* 26954 27053: gap of 100 bp
* 27054 27756: contig of 703 bp in length
* 27757 27856: gap of 100 bp
* 27857 28554: contig of 698 bp in length
* 28555 28654: gap of 100 bp
* 28655 29348: contig of 694 bp in length
* 29349 29448: gap of 100 bp
* 29449 30148: contig of 700 bp in length
* 30149 30248: gap of 100 bp
* 30249 30953: contig of 705 bp in length
* 30954 31053: gap of 100 bp
* 31054 31742: contig of 689 bp in length
* 31743 31842: gap of 100 bp
* 31843 32547: contig of 705 bp in length
* 32548 32647: gap of 100 bp
* 32648 33352: contig of 705 bp in length
* 33353 33452: gap of 100 bp
* 33453 34146: contig of 694 bp in length
* 34147 34246: gap of 100 bp
* 34247 34952: contig of 706 bp in length
* 34953 35052: gap of 100 bp
* 35053 35753: contig of 701 bp in length
* 35754 35853: gap of 100 bp
* 35854 36537: contig of 684 bp in length
* 36538 36637: gap of 100 bp
* 36638 37322: contig of 685 bp in length
* 37323 37422: gap of 100 bp
* 37423 38115: contig of 693 bp in length
* 38116 38215: gap of 100 bp
* 38216 38917: contig of 702 bp in length
* 38918 39017: gap of 100 bp
* 39018 39723: contig of 706 bp in length
* 39724 39823: gap of 100 bp
* 39824 40526: contig of 703 bp in length
* 40527 40626: gap of 100 bp
* 40627 41323: contig of 697 bp in length
* 41324 41423: gap of 100 bp
* 41424 42138: contig of 715 bp in length
* 42139 42238: gap of 100 bp
* 42239 42946: contig of 708 bp in length
* 42947 43046: gap of 100 bp
* 43047 43766: contig of 720 bp in length
* 43767 43866: gap of 100 bp
* 43867 44567: contig of 701 bp in length

```

```

Query Match      74.2%; Score 17.8; DB 2; Length 69320;
Best Local Similarity 90.5%; Pred. No. 97;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

```

QY 4 AGTGTGTCGAGGCATCTAGT 24
|||||
Db 56929 AGTGTGTCGAGGCATAGT 56909

```

```

RESULT 12
AC048373/c
LOCUS
DEFINITION Homo sapiens chromosome 11 clone RP11-139C6 map 11, WORKING DRAFT
AC048373
ACCESSION AC048373
VERSION AC048373.2 GI:7798808
KEYWORDS HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens.

```

## ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 152631)

Birren,B., Linton,L., Nusbaum,C. and Lander,E.

Homo sapiens chromosome 11, clone RP11-139C6

Unpublished

2 (bases 1 to 152631)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,  
Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,

Boguslavsky,L., Boukhgalter,B., Brown,A., Burkett,G.,

Campomione,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,

Collymore,A., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S.,

Dodge,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,

Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,

Grand-Pierre,N., Grant,G., Hagos,B., Hearford,A., Horton,L.,

Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,

Klein,J., LaRocque,K., Lamazares,R., Landers,T., Lehoczy,J.,

Levine,R., Liu,G., Locke,K., Macdonald,P., Marquis,N.,

McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,

Meldrim,J., Meneus,L., Mihova,T., Miranda,C., Miensga,V., Morrow,J.,

Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,

O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,

Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,

Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,

Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,

Tessfaye,S., Theodore,J., Tirrell,A., Travers,M., Trigglio,J.,

Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,

Young,G., Zainoun,J., Zimmer,A. and Zody,M.

Direct Submission

Submitted (14-APR-2000) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

On May 13, 2000 this sequence version replaced gi:7549705.

All repeats were identified using RepeatMasker:

Smit, A.F.A. &amp; Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L8701

Center clone name: 139\_C\_6

----- Summary Statistics

Sequencing vector: M13; M77815; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 145913 bases at least Q40

Consensus quality: 149504 bases at least Q30

Consensus quality: 150798 bases at least Q20

Insert size: 151000; agarose-1p

Insert size: 151631; sum-of-contigs

Quality coverage: 4.8 in Q20 bases; agarose-1p

Quality coverage: 4.8 in Q20 bases; sum-of-contigs

-----

\* NOTE: This is a 'working draft' sequence. It currently

\* consists of 11 contigs. The true order of the pieces

\* is not known and their order in this sequence record is

\* arbitrary. Gaps between the contigs are represented as

\* runs of N, but the exact sizes of the gaps are unknown.

\* This record will be updated with the finished sequence

\* as soon as it is available and the accession number will

\* be preserved.

\* 1 1150: contig of 1150 bp in length

\* 1151 1250: gap of 100 bp

\* 1251 6488: contig of 5238 bp in length

\* 6489 6589: gap of 100 bp

\* 6589 11741: contig of 5153 bp in length

\* 11742 11841: gap of 100 bp

\* 11842 16228: contig of 4387 bp in length

\* 16229 16328: gap of 100 bp

\* 16329 28016: contig of 11688 bp in length

```

* 28017 28116: gap of 100 bp
* 28117 45296: contig of 17180 bp in length
* 45297 45396: gap of 100 bp
* 45397 59567: contig of 14171 bp in length
* 59568 59667: gap of 100 bp
* 59668 76149: contig of 16482 bp in length
* 76150 76249: gap of 100 bp
* 76250 92365: contig of 16116 bp in length
* 92366 92465: gap of 100 bp
* 92466 119998: contig of 27533 bp in length
* 119999 120098: gap of 100 bp
* 120099 152631: contig of 32533 bp in length.

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## FEATURES

## source

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Location/Qualifiers
1. .152631
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/db_xref="taxon:9606"
/chromosome="11"
/map="11"
/clone="RP11-139C6"
/clone_lib="RPC1-11 Human Male BAC"
1. .1150
/feature="assembly_fragment"
1251. 6488
/feature="assembly_fragment"
6589. 111741
/feature="assembly_fragment"
11842. 16228
/feature="assembly_fragment"
clone_end.SP6
vector_side:left
16329. 28016
/feature="assembly_fragment"
clone_end.T7
vector_side:right
28117. 45296
/feature="assembly_fragment"
45397. 59567
/feature="assembly_fragment"
59668. 76149
/feature="assembly_fragment"
76250. 92365
/feature="assembly_fragment"
92466. 119998
/feature="assembly_fragment"
120099. 152631
/feature="assembly_fragment"

```

```

BASE COUNT 49516 a 26157 c 27027 g 48926 t 1005 others
ORIGIN

```

## misc\_feature

## misc\_feature

## misc\_feature

## misc\_feature

## misc\_feature

## misc\_feature

## misc\_feature

## misc\_feature

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## misc\_feature

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## misc\_feature

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## misc\_feature

## misc\_feature

## misc\_feature

REFERENCE  
AUTHORS

2 (bases 1 to 156507)  
 Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,  
 Anderson,S., Baldwin,J., Barna,N., Beckerly,R., Bada,F.,  
 Boguslavsky,L., Boukhgalter,B., Brown,A., Burkett,G., Castle,A.,  
 Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,  
 DeArelano,K., Dewar,K., Domino,M., Doyle,M., Fenestor,J.,  
 Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J.,  
 Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,  
 Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,  
 Landers,T., Lehoczy,J., Levine,R., Lieu,C., Liu,S., Locke,K.,  
 Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,  
 McPheeters,R., McElroy,J., Meneses,L., Morrow,J., Naylor,J.,  
 Norman,C.H., O'Connor,T., O'Donnell,P., Olivari,T.M., Peterson,K.,  
 Pierre,N., Pisani,C., Pollara,V., Raymond,C., Riley,R., Rothman,D.,  
 Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,  
 Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,  
 Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,  
 Zimmer,A. and Zody,M.

TITLE  
JOURNAL

Direct Submission  
 Submitted (22-JAN-2000) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA

REFERENCE  
AUTHORS

3 (bases 1 to 156507)  
 Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,  
 Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B.,  
 Brown,A., Camarata,J., Campiano,A., Chang,J., Chazaro,B.,  
 Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,  
 Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,  
 Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,  
 Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,  
 Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,  
 Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,  
 Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,  
 Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,  
 McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrim,J.,  
 Meneses,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,  
 Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D.,  
 Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,  
 Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,  
 Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupbach,R.,  
 Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,  
 Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,  
 Topham,K., Travers,M., Travis,N., Triggillo,J., Vassiliev,H.,  
 Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,  
 Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

TITLE  
JOURNAL

Direct Submission  
 Submitted (03-NOV-2001) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA  
 On Oct 21, 2001 this sequence version replaced gi:16259015.

## COMMENT

All repeats were identified using RepeatMasker:  
 Smit, A.F.A. & Green, P. (1996-1997)  
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L4204

Center clone name: 68\_C8

FEATURES  
source

```

Location/Qualifiers
1. .156507
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="11"
/map="11"
/clone="RP11-68C8"
/clone_lib="RPC1-11 Human Male BAC"
complement(132..1126)
/rpt_family="LIMC3"
repeat_region 1129..1388
/rpt_family="LFR26B"
repeat_region 1433..1588

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## RESULT 13

## AC021914

LOCUS AC021914 156507 bp DNA linear PRI 03-NOV-2001  
 DEFINITION Homo sapiens chromosome 11, clone RP11-68C8, complete sequence.

## ACCESSION

## AC021914

## VERSION

## AC021914.7

## KEYWORDS

## HTG.

## SOURCE

## Homo sapiens.

## ORGANISM

## Homo sapiens

## Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

## Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

## REFERENCE

## 1 (bases 1 to 156507)

## AUTHORS

## Birren,B., Linton,L., Nusbaum,C. and Lander,E.

## TITLE

## Homo sapiens chromosome 11, clone RP11-68C8

## JOURNAL

## Unpublished



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repeat_region /rpt_family="LTR67"
repeat_region complement(1794..2350)
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repeat_region 3141..3440
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repeat_region 5555..5749
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repeat_region 5896..6215
repeat_region /rpt_family="AluSc"
repeat_region 6683..6721
repeat_region /rpt_family="AT_rich"
repeat_region complement(6896..8100)
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repeat_region /rpt_family="MERS2A"
repeat_region complement(9688..10187)
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repeat_region complement(10188..10601)
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repeat_region complement(11102..11204)
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repeat_region 11218..11462
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repeat_region 11553..11644
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repeat_region 16721..16864
repeat_region /rpt_family="FLAM_C"
repeat_region complement(16867..17095)
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repeat_region 17119..17522
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repeat_region complement(17896..17983)
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repeat_region 18245..18266
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repeat_region 18587..18612
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repeat_region 18976..19004
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repeat_region complement(20451..20507)
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repeat_region 20944..20985
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repeat_region complement(22094..22386)
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repeat_region 22673..22709
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Query Match 74.2%; Score 17.8; DB 9; Length 156507;  
 Best Local Similarity 90.5%; Pred. No. 94;  
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCTAGTCTGTCGAGCATCT 21  
 ||||| ||||| ||||| |||||  
 Db 57570 TCTAGTCTGTCGAGCATCT 57590

RESULT 14  
 AP002378/c

LOCUS Homo sapiens genomic DNA, chromosome 11q, clone:RP11-351D3,  
 DEFINITION complete sequences.  
 ACCESSION AP002378  
 VERSION AP002378.3 GI:13810520  
 KEYWORDS HTG.

SOURCE Homo sapiens DNA, clone:RP11-351D3.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE 1 (sites)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
 Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.

TITLE Homo sapiens genomic DNA  
 JOURNAL Published Only in Database (2000)

REFERENCE 2 (bases 1 to 177429)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
 Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.  
 TITLE Direct Submission  
 JOURNAL Submitted (29-MAY-2000) Masahira Hattori, The Institute of Physical

and Chemical Research (RIKEN), Genomic Sciences Center (GSC);  
1-7-22 Suehiro-chou, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
(E-mail: hattori@gsc.riken.go.jp, URL: http://hgp.gsc.riken.go.jp/,  
Tel: 81-45-503-9111, Fax: 81-45-503-9170)

On Apr 26, 2001 this sequence version replaced gi:11136462.

#### FEATURES

source

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Location/Qualifiers

BASE COUNT 57127 a 30586 c 31846 g 57870 t  
ORIGIN

Query Match 74.2%; Score 17.8; DB 9; Length 177429;

Best Local Similarity 90.5%; Pred. No. 94;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCTAGTGTGTCGAGGCATCT 21

||||||| |||||||||

Db 102035 TCTAGTGGAGTCGAGGCATCT 102015

#### RESULT 15

AP000760/c

LOCUS

DEFINITION

AP000760

complete sequences.

AP000760.4 GI:13810525

HTG.

AP000760

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

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/db\_xref="taxon:9606"

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/clone="RP11-779L16"

Location/Qualifiers

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ORIGIN

Query Match 74.2%; Score 17.8; DB 9; Length 177488;

Best Local Similarity 90.5%; Pred. No. 94;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCTAGTGTGTCGAGGCATCT 21

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Db 3629 TCTAGTGGAGTCGAGGCATCT 3609

Search completed: January 3, 2003, 23:57:17

Job time : 371.151 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:02 ; Search time 40.1262 Seconds  
(without alignments)  
1346.950 Million cell updates/sec

Title: US-09-787-562-11

Perfect score: 24

Sequence: 1 ttctagtctgtgcaggcatctagt 24

Scoring table: IDENTITY\_NUC

Gapop 10\_0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	24	100.0	24	21	AAA12003
2	24	100.0	237	21	AAA12001
3	17.8	74.2	2817	18	AAV74606
C 4	17.4	72.5	72	20	AAZ11441
C 5	17.4	72.5	72	21	AAA11998
C 6	17.4	72.5	72	21	AAA12024
C 7	17.4	72.5	72	21	AAA12025
C 8	16.6	69.2	39	8	AAV70749
C 9	16.6	69.2	111	22	AAV25330
					Murine HIF-1 space
					Murine PGK HRE der
					Staphylococcus aur
					Mutant HRE-contain
					Murine PGK HRE-con
					Murine PGK HRE HIF
					Murine PGK HRE-con
					Sequence of portio
					Human breast cance

C 10	16.6	69.2	115	22	AAV16487	Human breast cance
C 11	16.6	69.2	192	23	ABV14349	Human prostate exp
C 12	16.6	69.2	202	21	AAV09061	Human prostate pro
C 13	16.6	69.2	450	23	ABV44262	Human prostate exp
C 14	16.6	69.2	479	23	ABV35433	Human prostate exp
C 15	16.6	69.2	726	22	AAV10807	Human Janus kinase
C 16	16.6	69.2	749	22	ABA09516	Human endozepine-r
C 17	16.6	69.2	749	22	AAV99188	Human protein enco
C 18	16.6	69.2	1498	22	AAV78096	Human immune/haema
C 19	16.6	69.2	1593	22	AAV81568	Human endozepine-1
C 20	16.6	69.2	1677	24	AAV43759	Human NOV2 gene se
C 21	16.6	69.2	1727	21	AAA47432	Sequence encoding
C 22	16.6	69.2	1747	24	ABN88660	Human NOV1 encodin
C 23	16.6	69.2	1930	22	AAV84889	Human SEC8 nucleic
C 24	16.6	69.2	2056	22	AAV01220	DNA encoding human
C 25	16.6	69.2	2056	22	AAV84893	Human SEC12 nucle
C 26	16.6	69.2	2510	22	AAV51825	Human polynucleoti
C 27	16.6	69.2	3807	17	AAV30862	Protein tyrosine k
C 28	16.6	69.2	3887	24	AAV43754	Human NOV1a gene s
C 29	16.6	69.2	3920	24	AAV43755	Human NOV1b gene s
C 30	16.6	69.2	3920	24	AAV43756	Human NOV1c gene s
C 31	16.6	69.2	3920	24	AAV43757	Human NOV1d gene s
C 32	16.6	69.2	3920	24	AAV43758	Human NOV1e gene s
C 33	16.6	69.2	6781	19	AAV45824	Maize phosphenoip
C 34	16.6	69.2	7559	20	AAV82011	Maize phosphenoip
C 35	16.6	69.2	26059	22	AAV69104	Human immune/haema
C 36	16.2	67.5	557	24	ABV77504	Bacillus clausil g
C 37	16.2	67.5	902	18	AAV75089	Staphylococcus aur
C 38	16.2	67.5	1659	20	AAV27698	HSP47 gene fragmen
C 39	16.2	67.5	1698	21	AAV81571	N. meningitidis pa
C 40	16.6	66.7	72	20	AAV11440	HRE-containing enh
C 41	16.6	66.7	72	21	AAV12023	Murine PGK HRE HIF
C 42	16.6	66.7	80	20	AAV11431	HRE element contai
C 43	16.6	66.7	100	21	AAV12060	EIAV U3 enhancer r
C 44	16.6	66.7	114	21	AAV12061	EIAV U3 enhancer r
C 45	16.6	66.7	159	22	AAV71452	Human foetal liver

#### ALIGNMENTS

RESULT 1

AAA12003

ID AAA12003 standard; DNA: 24 BP.

XX

XX

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XX

PT Novel polynucleotide constructs comprising at least two repeats of a  
 PT hypoxia response element useful for driving expression of nucleic acids  
 PT of interest in a cell under hypoxic conditions -  
 XX  
 PS Disclosure; Page 10; 155pp; English.  
 XX  
 CC This invention describes novel polynucleotide comprising at least 2  
 CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible  
 CC factor (HIF) consensus binding sites within each of the 2 repeats are  
 CC separated by a spacer of at least 20 contiguous nucleotides. The products  
 CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic  
 CC activity and can be used for gene therapy. The polynucleotides are useful  
 CC for delivering nucleic acids of interest to mammalian cells. Lentiviral  
 CC vectors are responsive to hypoxic agents and to agents that mimic  
 CC hypoxia. This regulation can be harnessed in vitro to enhance the  
 CC production of the vector and can be used in vivo to regulate gene  
 CC expression in response to a physiological signal. The vectors have  
 CC utility in disease, where ischaemia, including hypoxia, is a feature,  
 CC e.g. cardiovascular disease, peripheral arterial disease, cancer and  
 CC arthritis. The novel regulatory construct is capable of driving very high  
 CC levels of transcription under conditions of hypoxia whilst providing only  
 CC low basal levels of transcription under normal oxygen conditions. The  
 CC polynucleotide construct targets cells within a tumor mass that are under  
 CC conditions of hypoxia without affecting normal surrounding tissue. This  
 CC sequence represents a murine HIF-1 DNA spacer which is used in the method  
 CC of the invention.  
 XX  
 SQ Sequence 24 BP; 4 A; 5 C; 7 G; 8 T; 0 other;

Query Match 100.0%; Score 24; DB 21; Length 24;  
 Best Local Similarity 100.0%; Pred. No. 0.013;  
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 TCTAGTGTCTGTCAGGCATCTAGT 24  
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 Db 1 TCTAGTGTCTGTCAGGCATCTAGT 24

RESULT 2  
 AAAL2001  
 ID AAAL2001 standard; DNA; 237 BP.  
 AC AAAL2001;  
 XX  
 DT 14-AUG-2000 (first entry)  
 XX  
 DE Murine PGK HRE derived promoter OBHrel1 DNA.  
 XX  
 KW HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;  
 KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;  
 KW cardiovascular disease; peripheral arterial disease; cancer;  
 KW phosphoglycerate kinase; PGK; murine; promoter; OBHrel1; ds.  
 XX  
 OS Mus sp.  
 XX  
 PN WO200017371-A1.  
 XX  
 PD 30-MAR-2000.  
 XX  
 PF 22-SEP-1999; 99WO-GB03181.  
 XX  
 XX 23-SEP-1998; 98WO-GB02885.  
 PR 28-JAN-1999; 99GB-0001906.  
 PR 16-FEB-1999; 99GB-0003538.  
 XX  
 XX (OXFO-) OXFORD BIOMEDICA UK LTD.  
 PA  
 XX  
 PI Binley KM, Naylor S;  
 XX  
 DR WPI; 2000-283595/24.  
 XX  
 XX Novel polynucleotide constructs comprising at least two repeats of a  
 PT hypoxia response element useful for driving expression of nucleic acids

PT of interest in a cell under hypoxic conditions -  
 XX  
 PS Example 1; Page 68; 155pp; English.  
 XX  
 CC This invention describes novel polynucleotide comprising at least 2  
 CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible  
 CC factor (HIF) consensus binding sites within each of the 2 repeats are  
 CC separated by a spacer of at least 20 contiguous nucleotides. The products  
 CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic  
 CC activity and can be used for gene therapy. The polynucleotides are useful  
 CC for delivering nucleic acids of interest to mammalian cells. Lentiviral  
 CC vectors are responsive to hypoxic agents and to agents that mimic  
 CC hypoxia. This regulation can be harnessed in vitro to enhance the  
 CC production of the vector and can be used in vivo to regulate gene  
 CC expression in response to a physiological signal. The vectors have  
 CC utility in disease, where ischaemia, including hypoxia, is a feature,  
 CC e.g. cardiovascular disease, peripheral arterial disease, cancer and  
 CC arthritis. The novel regulatory construct is capable of driving very high  
 CC levels of transcription under conditions of hypoxia whilst providing only  
 CC low basal levels of transcription under normal oxygen conditions. The  
 CC polynucleotide construct targets cells within a tumor mass that are under  
 CC conditions of hypoxia without affecting normal surrounding tissue. This  
 CC sequence represents a murine phosphoglycerate kinase (PGK) HRE derived  
 CC promoter OBHrel1 which is described in the method of the invention.  
 XX  
 SQ Sequence 237 BP; 43 A; 82 C; 56 G; 56 T; 0 other;

Query Match 100.0%; Score 24; DB 21; Length 237;  
 Best Local Similarity 100.0%; Pred. No. 0.019;  
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 TCTAGTGTCTGTCAGGCATCTAGT 24  
 |||||  
 Db 25 TCTAGTGTCTGTCAGGCATCTAGT 48

RESULT 3  
 AAV74606  
 ID AAV74606 standard; DNA; 2817 BP.  
 AC AAV74606;  
 XX  
 DT 16-MAR-1999 (first entry)  
 XX  
 DE Staphylococcus aureus contig SEQ ID #295.  
 XX  
 KW Computer readable medium; vaccine; S aureus infection; immunodetection;  
 KW cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;  
 KW skin infection; surgical wound infection; scalded skin syndrome;  
 KW toxic shock syndrome; ds.  
 XX  
 OS Staphylococcus aureus.  
 XX  
 PH Key Location/Qualifiers  
 FT misc\_feature 601..860  
 FT /\*tag= a  
 FT /note= "these bases represent a line of missing text in  
 FT the sequence listing in the specification. They  
 FT are included to maintain the nucleotide numbering  
 FT given in the specification for this DNA sequence"  
 FT misc\_feature 2401..2460  
 FT /\*tag= b  
 FT /note= "these bases represent a line of missing text in  
 FT the sequence listing in the specification. They  
 FT are included to maintain the nucleotide numbering  
 FT given in the specification for this DNA sequence"  
 FT  
 XX EP786519-A2.  
 PN  
 XX 30-JUL-1997.  
 PD  
 XX 07-JAN-1997; 97EP-0100117.  
 PF  
 XX

```

PR 05-JAN-1996; 96US-0009861.
XX (HUMA-) HUMAN GENOME SCI INC.
PA Barash SC, Choi GH, Dillon PJ, Fannon MR, Kunsch CA;
PI Rosen CA;
XX WPI; 1997-374922/35.
XX Polynucleotide(s) and proteins derived from Staphylococcus aureus
PT stored on computer readable medium and used in the production of
PT anti-S.aureus vaccines
XX Claim 1; Page 1120-1122; 3271pp; English.
XX This sequence represents one of 5191 Staphylococcus aureus DNA sequences
CC of the invention. The DNA sequences are recorded on a computer readable
CC medium, preferably selected from a floppy or hard disk, random access
CC memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
CC the S.aureus DNA sequences allows putative functions to be assigned so
CC that protein-encoding or regulatory regions of commercial, therapeutic or
CC industrial importance can be obtained. Specifically, sequences which are
CC likely to encode antigens have been identified and these polypeptides can
CC be used in a vaccine composition against S.aureus infection. The
CC polypeptides can also be used in a kit for the immunodetection of
CC S.aureus in a sample. S.aureus is implicated in numerous human diseases,
CC including cellulitis, eyelid infections, food poisoning, osteomyelitis,
CC skin and surgical wound infections, scalded skin syndrome, toxic shock
CC syndrome, etc. Organisms transformed with the DNA sequences can be used
CC for recombinant production of the polypeptides. The new DNA sequences
CC (and their fragments) are useful as primers or probes for isolating
CC homologues of any of the S.aureus DNA sequences contained on the
CC computer readable medium.
XX
SQ Sequence 2817 BP; 912 A; 330 C; 446 G; 999 T; 130 other;
Query Match 74.2%; Score 17.8; DB 18; Length 2817;
Best Local Similarity 90.5%; Pred. No. 36;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 4 AGTGTGCGTCAGGCATCTAGT 24
DB 1105 ATTGTGTGCGAGGCATCTAGT 1125
RESULT 4
AAZ11441/C
ID AAZ11441 standard; DNA; 72 BP.
AC AAZ11441;
XX
XX 26-OCT-1999 (first entry)
XX Mutant HRE-containing enhancer MUT PGK18+++
XX
XX Retroviral vector; functional splice donor site; hybrid viral vector;
KW functional splice acceptor site; in vivo gene delivery; therapeutic;
KW lentiviral vector; modified hematopoietic stem cell; MHC; tumour;
KW ischemia; hypoxia response element; HRE; hypoxia; promoter; ds.
XX Synthetic.
OS Mus sp.
XX
XX WO9915684-A2.
XX
XX 01-APR-1999.
XX
XX 23-SEP-1998; 98WO-GB02885.
XX
XX 25-SEP-1997; 97GB-0020465.
XX
XX 23-SEP-1997; 97GB-0020216.
XX
XX (OXFO-) OXFORD BIOMEDICA UK LTD.
XX
XX Bebbington C, Binley KM, Lewis C, Naylor S;
XX WPI; 1999-263482/22.
XX New retroviral vectors, for, e.g. delivering nucleotide sequences to
XX solid tumor sites
XX Example 5 (page 77); Fig 13 (Page 16/43); 288pp; English.
XX The invention relates to a retroviral vector (RVV) comprising a
CC functional splice donor site (FSDS) and a functional splice acceptor site
CC (FSAS) where: (i) the FSDS and the FSAS flank a first nucleotide sequence
CC of interest (NOI); (ii) the FSDS is upstream of the FSAS; (iii) the RVV
CC is derived from a retroviral pro-vector; (iv) the retroviral pro-vector
CC comprises a first nucleotide sequence (NS) capable of yielding the FSDS
CC and a second NS capable of yielding the FSAS; and (v) the first NS is
CC downstream of the second NS, such that the RVV is formed as a result of
CC reverse transcription of the retroviral pro-vector. A hybrid viral vector
CC (VV) system for in vivo gene delivery, which system comprises a primary
CC a first target cell and of expressing the secondary target cell, where the
CC vector is capable of transducing a secondary target cell, where the
CC primary vector is obtainable from or is based on a adenoviral vector and
CC the secondary VV is obtainable from or is based on a RVV preferably a
CC delivering NOIs to one or more target sites. The systems can be used for
CC therapeutic or diagnostic agents. The methods are used particularly for
CC producing modified hematopoietic stem cells (MHSCs) to deliver NOIs to
CC sites such as solid tumours which are characterised by ischemia, such as
CC hypoxia or low glucose concentration. The system permits the stable
CC expression of NOIs in targeted cells, e.g. rapidly dividing cells. The
CC present sequence represents a mutant HRE-containing enhancer sequence
CC derived from PGK gene.
XX
SQ Sequence 72 BP; 16 A; 25 C; 12 G; 19 T; 0 other;
Query Match 72.5%; Score 17.4; DB 20; Length 72;
Best Local Similarity 94.7%; Pred. No. 34;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TCTAGTGTCTGTCGAGGCAT 19
DB 29 TCTAGTGTCTGTCGAGGCAT 11
RESULT 5
AAZ11998/C
ID AAZ11998 standard; DNA; 72 BP.
AC AAZ11998;
XX
XX 14-AUG-2000 (first entry)
XX
XX Murine PGK HRE-containing enhancer element WT PGK18+++ DNA.
XX HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;
KW cardiant; cytostatic; antiarthritic; gene therapy; ischemia; arthritis;
KW cardiovascular disease; peripheral arterial disease; cancer;
KW phosphoglycerate kinase; PGK; murine; enhancer; ds.
XX
XX Mus sp.
XX
XX WO200017371-A1.
XX
XX 30-MAR-2000.
XX
XX 22-SEP-1999; 99WO-GB03181.
XX
XX 23-SEP-1998; 98WO-GB02885.
XX
XX 28-JAN-1999; 99GB-0001906.
XX
XX 16-FEB-1999; 99GB-0003538.
XX

```

PA (OXFO-) OXFORD BIOMEDICA UK LTD.  
XX  
XX  
PI Binley KM, Naylor S;  
XX  
XX WPT: 2000-283595/24.  
XX  
XX Novel polynucleotide constructs comprising at least two repeats of a  
PT hypoxia response element useful for driving expression of nucleic acids  
PT of interest in a cell under hypoxic conditions -  
XX  
XX Example 6; Page 79; 155pp; English.  
XX  
XX This invention describes novel polynucleotide comprising at least 2  
XX repeats of a hypoxia response element (HRE), where the hypoxia-inducible  
CC factor (HIF) consensus binding sites within each of the 2 repeats are  
CC separated by a spacer of at least 20 contiguous nucleotides. The products  
CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic  
CC activity and can be used for gene therapy. The polynucleotides are useful  
CC for delivering nucleic acids of interest to mammalian cells. Lentiviral  
CC vectors are responsive to hypoxic agents and to agents that mimic  
CC hypoxia. This regulation can be harnessed in vitro to enhance the  
CC production of the vector and can be used in vivo to regulate gene  
CC expression in response to a physiological signal. The vectors have  
CC utility in disease, where ischaemia, including hypoxia, is a feature,  
CC e.g. cardiovascular disease, peripheral arterial disease, cancer and  
CC arthritis. The novel regulatory construct is capable of driving very high  
CC levels of transcription under conditions of hypoxia whilst providing only  
CC low basal levels of transcription under normal oxygen conditions. The  
CC polynucleotide construct targets cells within a tumor mass that are under  
CC conditions of hypoxia without affecting normal surrounding tissue. This  
CC sequence represents a murine phosphoglycerate kinase (PGK) HRE derived  
CC enhancer which is described in the method of the invention.  
XX  
XX Sequence 72 BP; 16 A; 25 C; 12 G; 19 T; 0 other;  
SQ  
Query Match 72.5%; Score 17.4; DB 21; Length 72;  
Best Local Similarity 94.7%; Pred. No. 34;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 1 TCTAGTGTCTGTCAGGCAT 19  
|||||  
Db 29 TCTAGTGTCTGTCAGGAAT 11  
RESULT 6  
AAAL2024/c  
ID AAAL2024 standard; DNA; 72 BP.  
XX  
XX AC AAAL2024;  
XX  
XX DT 14-AUG-2000 (first entry)  
XX  
XX DE Murine PGK HRE HIF-1 binding site mutant DNA.  
XX  
XX HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;  
KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;  
KW cardiovascular disease; peripheral arterial disease; cancer;  
KW phosphoglycerate kinase; PGK; murine; ds.  
XX  
XX OS Mus sp.  
XX  
XX PN WO200017371-A1.  
XX  
XX PD 30-MAR-2000.  
XX  
XX PF 22-SEP-1999; 99WO-GB03181.  
XX  
XX PR 23-SEP-1998; 98WO-GB02885.  
XX  
XX PR 28-JAN-1999; 99GB-0001906.  
XX  
XX PR 16-FEB-1999; 99GB-0003538.  
XX  
XX PA (OXFO-) OXFORD BIOMEDICA UK LTD.  
XX  
XX PI Binley KM, Naylor S;  
XX

PI Binley KM, Naylor S;  
XX  
XX WPI: 2000-283595/24.  
XX  
XX Novel polynucleotide constructs comprising at least two repeats of a  
PT hypoxia response element useful for driving expression of nucleic acids  
PT of interest in a cell under hypoxic conditions -  
XX  
XX Example 4; Page 76; 155pp; English.  
XX  
XX This invention describes novel polynucleotide comprising at least 2  
XX repeats of a hypoxia response element (HRE), where the hypoxia-inducible  
CC factor (HIF) consensus binding sites within each of the 2 repeats are  
CC separated by a spacer of at least 20 contiguous nucleotides. The products  
CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic  
CC activity and can be used for gene therapy. The polynucleotides are useful  
CC for delivering nucleic acids of interest to mammalian cells. Lentiviral  
CC vectors are responsive to hypoxic agents and to agents that mimic  
CC hypoxia. This regulation can be harnessed in vitro to enhance the  
CC production of the vector and can be used in vivo to regulate gene  
CC expression in response to a physiological signal. The vectors have  
CC utility in disease, where ischaemia, including hypoxia, is a feature,  
CC e.g. cardiovascular disease, peripheral arterial disease, cancer and  
CC arthritis. The novel regulatory construct is capable of driving very high  
CC levels of transcription under conditions of hypoxia whilst providing only  
CC low basal levels of transcription under normal oxygen conditions. The  
CC polynucleotide construct targets cells within a tumor mass that are under  
CC conditions of hypoxia without affecting normal surrounding tissue. This  
CC sequence represents a murine wild type phosphoglycerate kinase (PGK) HRE  
CC HIF-1 binding site which is described in the method of the invention.  
XX  
XX Sequence 72 BP; 16 A; 25 C; 12 G; 19 T; 0 other;  
SQ  
Query Match 72.5%; Score 17.4; DB 21; Length 72;  
Best Local Similarity 94.7%; Pred. No. 34;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 1 TCTAGTGTCTGTCAGGCAT 19  
|||||  
Db 29 TCTAGTGTCTGTCAGGAAT 11  
RESULT 7  
AAAL2025/c  
ID AAAL2025 standard; DNA; 72 BP.  
XX  
XX AC AAAL2025;  
XX  
XX DT 14-AUG-2000 (first entry)  
XX  
XX DE Murine PGK HRE-containing enhancer element MT PGK18+++ DNA.  
XX  
XX HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;  
KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;  
KW cardiovascular disease; peripheral arterial disease; cancer;  
KW phosphoglycerate kinase; PGK; murine; enhancer; ds.  
XX  
XX OS Mus sp.  
XX  
XX PN WO200017371-A1.  
XX  
XX PD 30-MAR-2000.  
XX  
XX PF 22-SEP-1999; 99WO-GB03181.  
XX  
XX PR 23-SEP-1998; 98WO-GB02885.  
XX  
XX PR 28-JAN-1999; 99GB-0001906.  
XX  
XX PR 16-FEB-1999; 99GB-0003538.  
XX  
XX PA (OXFO-) OXFORD BIOMEDICA UK LTD.  
XX  
XX PI Binley KM, Naylor S;  
XX

DR WPI; 2000-283595/24.  
 XX  
 PT Novel polynucleotide constructs comprising at least two repeats of a  
 PT hypoxia response element useful for driving expression of nucleic acids  
 PT of interest in a cell under hypoxic conditions  
 PS Example 6; Page 79; 155pp; English.  
 XX  
 CC This invention describes novel polynucleotide comprising at least 2  
 CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible  
 CC factor (HIF) consensus binding sites within each of the 2 repeats are  
 CC separated by a spacer of at least 20 contiguous nucleotides. The products  
 CC of the invention have vasotropic, cardiac, cytostatic and antiarthritic  
 CC activity and can be used for gene therapy. The polynucleotides are useful  
 CC for delivering nucleic acids of interest to mammalian cells. Lentiviral  
 CC vectors are responsive to hypoxic agents and to agents that mimic  
 CC hypoxia. This regulation can be harnessed in vitro to enhance the  
 CC production of the vector and can be used in vivo to regulate the  
 CC expression in response to a physiological signal. The vectors have  
 CC utility in disease, where ischemia, including hypoxia, is a feature,  
 CC e.g. cardiovascular disease, peripheral arterial disease, cancer and  
 CC arthritis. The novel regulatory construct is capable of driving very high  
 CC levels of transcription under conditions of hypoxia whilst providing only  
 CC low basal levels of transcription under normal oxygen conditions. The  
 CC polynucleotide construct targets cells within a tumor mass that are under  
 CC conditions of hypoxia without affecting normal surrounding tissue. This  
 CC sequence represents a murine phosphoglycerate kinase (PGK) HRE derived  
 CC enhancer which is described in the method of the invention.  
 XX  
 SQ Sequence 72 BP; 16 A; 25 C; 12 G; 19 T; 0 other;  
 Query Match 72.5%; Score 17.4; DB 21; Length 72;  
 Best Local Similarity 94.7%; Pred. No. 34;  
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 TCTAGTGTCTGTCGAGGCAT 19  
 Db 29 TCTAGTGTCTGTCGAGGCAT 11  
 RESULT 8  
 ID AAN70749/C  
 XX AAN70749 standard; DNA; 39 BP.  
 AC AAN70749;  
 XX  
 DT 03-OCT-2002 (updated)  
 DT 04-FEB-1991 (first entry)  
 XX  
 DE Sequence of portion of Drosophila heat-shock hybrid gene on p17-lys  
 DE showing heat shock protein (Hsp70) promoter.  
 XX  
 KW Gene expression; transcription control element;  
 KW Drosophila 70kD heat shock protein promoter; ss;  
 KW chicken lysozyme gene; human hsp70 gene.  
 XX  
 OS Drosophila melanogaster.  
 XX  
 FH Key Location/Qualifiers  
 FT misc\_feature 1..14  
 FT /tag= a  
 FT /note="Hsp70 gene sequence"  
 XX  
 PN W08705935-A.  
 XX  
 XX 08-OCT-1987.  
 PD  
 PD 03-APR-1987; 87WO-US00805.  
 PF  
 XX 04-APR-1986; 86US-0848657.  
 PR  
 XX (BATT) BATTELLE MEMORIAL INST.  
 PA  
 XX

PI Bromley P, Voellmy R;  
 XX WPI; 1987-291648/41.  
 XX  
 PT DNA contg. structural gene and heat shock promoter - esp. from  
 PT Drosophila, allowing induction of protein expression in prokaryotic  
 PT or eukaryotic cells.  
 XX  
 PS Example; Fig 13; 105pp; English.  
 XX  
 CC The sequence in AAN70743 was cut from plasmid p51 and inserted into  
 CC pMC 1103 in front of its incomplete beta-galactosidase gene to form  
 CC recombinant plasmid pRV15 (specifically claimed) which was able to  
 CC express beta-galactosidase. pRV15 was digested with SmaI and SalI to  
 CC give a 7 kbp segment and this ligated with pSVod to form plasmid  
 CC 520. Separately, plasmid pR81 was digested with HindIII and NcoI.  
 CC The resulting fragments were ligated with a PvuII-NcoI fragment of  
 CC p520 to form the specifically claimed plasmid pR84 (AAN70744). The  
 CC heat shock promoters are used for the inducible expression of genes  
 CC encoding secretable and non-secretable proteins. A 1.5 kb fragment  
 CC from p522-lys and ligated to a 1.2 kb long fragment contg. a  
 CC functional heat shock promoter from p17 to give p17-lys.  
 CC (Updated on 03-OCT-2002 to add missing OS field.)  
 XX  
 SQ Sequence 39 BP; 12 A; 11 C; 10 G; 6 T; 0 other;  
 Query Match 69.2%; Score 16.6; DB 8; Length 39;  
 Best Local Similarity 82.6%; Pred. No. 78;  
 Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 2 CTAGTGTCTGTCGAGGCATCTAGT 24  
 Db 26 CCAGTGTCTGTCGAGGCATCTGT 4  
 RESULT 9  
 ID AAL25330/C  
 XX AAL25330 standard; cDNA; 111 BP.  
 AC AAL25330;  
 XX  
 DT 07-DEC-2001 (first entry)  
 XX  
 DE Human breast cancer expressed polynucleotide 17787.  
 DE  
 KW Human; breast cancer; cell marker; cytostatic; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN W0200151628-A2.  
 XX  
 PD 19-JUL-2001.  
 XX  
 PF 10-JAN-2001; 2001WO-US00798.  
 XX  
 XX 14-JAN-2000; 2000US-0176077.  
 PR 14-MAR-2000; 2000US-0189167.  
 PR 24-MAR-2000; 2000US-0192099.  
 PR 29-MAR-2000; 2000US-0193480.  
 PR 15-MAY-2000; 2000US-0205230.  
 PR 09-JUN-2000; 2000US-0211315.  
 PR 25-JUL-2000; 2000US-0220534.  
 XX  
 PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
 XX  
 PI Lillie J, Xu Y, Wang Y, Steinmann K;  
 XX  
 DR WPI; 2001-451856/48.  
 XX  
 PT New peptide useful as a marker for the diagnosis of breast cancer  
 XX  
 PS Claim 1; Page 3293; 3695pp; English.

XX The invention relates to human breast cancer expressed polynucleotides  
 CC (AAL07544-AAL26789) and methods of assessing whether a patient is  
 CC afflicted with breast cancer by examining the correlation between the  
 CC expression of certain markers and the cancerous state of breast cells.  
 CC The polynucleotides and encoded polypeptides are potential markers for  
 CC detecting, diagnosing, monitoring, characterising treating and  
 CC potentially preventing breast cancer. The polynucleotides and encoded  
 CC polypeptides are also useful for isolating compounds with cytostatic  
 CC activity.  
 XX  
 SQ Sequence 111 BP; 23 A; 30 C; 39 G; 19 T; 0 other;

Query Match 69.2%; Score 16.6; DB 22; Length 111;  
 Best Local Similarity 82.6%; Pred. No. 91;  
 Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTCTCGTCGAGGCATCTAGT 24  
 DB 105 CTAGTCTCGTCGAGGCATCTCGT 83

RESULT 10  
 AAL16487/c  
 ID AAL16487 standard; cDNA; 115 BP.  
 XX  
 AC AAL16487;  
 XX  
 XX 07-DEC-2001 (first entry)  
 XX  
 XX Human breast cancer expressed polynucleotide 8944.  
 XX  
 XX Human; breast cancer; cell marker; cytostatic; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 XX WO200151628-A2.  
 XX  
 XX 19-JUL-2001.  
 XX  
 XX 10-JAN-2001; 2001WO-US00798.

XX 14-JAN-2000; 2000US-0176077.  
 PR 14-MAR-2000; 2000US-0189167.  
 PR 24-MAR-2000; 2000US-0192099.  
 PR 29-MAR-2000; 2000US-0193480.  
 PR 15-MAY-2000; 2000US-0205230.  
 PR 09-JUN-2000; 2000US-0211315.  
 PR 25-JUL-2000; 2000US-0220534.  
 XX  
 PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
 XX  
 XX Lillie J, Xu Y, Wang Y, Steinmann K;  
 PI  
 XX WPI; 2001-451856/48.

XX New peptide useful as a marker for the diagnosis of breast cancer -  
 XX  
 PS Claim 1; Page 1618; 3695pp; English.

XX The invention relates to human breast cancer expressed polynucleotides  
 CC (AAL07544-AAL26789) and methods of assessing whether a patient is  
 CC afflicted with breast cancer by examining the correlation between the  
 CC expression of certain markers and the cancerous state of breast cells.  
 CC The polynucleotides and encoded polypeptides are potential markers for  
 CC detecting, diagnosing, monitoring, characterising treating and  
 CC potentially preventing breast cancer. The polynucleotides and encoded  
 CC polypeptides are also useful for isolating compounds with cytostatic  
 CC activity.  
 XX

SQ Sequence 115 BP; 24 A; 30 C; 41 G; 20 T; 0 other;

Query Match 69.2%; Score 16.6; DB 22; Length 115;

Best Local Similarity 82.6%; Pred. No. 91;  
 Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTCTCGTCGAGGCATCTAGT 24  
 DB 109 CTAGTCTCGTCGAGGCATCTCGT 87

RESULT 11  
 ABV14349/c  
 ID ABV14349 standard; cDNA; 192 BP.  
 XX  
 AC ABV14349;  
 XX  
 XX 13-SEP-2002 (first entry)  
 XX  
 XX Human prostate expression marker cDNA 14340.

XX Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
 KW pharmacogenomic marker; gene; ss.  
 XX  
 OS Homo sapiens.

XX  
 XX WO200160860-A2.  
 XX  
 XX 23-AUG-2001.  
 XX  
 XX 20-FEB-2001; 2001WO-US05171.

XX 17-FEB-2000; 2000US-183319P.  
 PR 16-MAR-2000; 2000US-189862P.  
 PR 25-MAY-2000; 2000US-207454P.  
 PR 09-JUN-2000; 2000US-211314P.  
 PR 18-JUL-2000; 2000US-219007P.  
 PR 13-DEC-2000; 2000US-255281P.

XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
 XX  
 XX Schlegel R, Endege WO, Monahan JE;  
 PI  
 XX WPI; 2001-662795/76.

XX Novel isolated nucleic acid molecule associated with cancerous state of  
 PT prostate cells and correlating with presence of prostate cancer, useful  
 PT for detecting presence of prostate cancer, stage of prostate cancer -  
 XX  
 PS Claim 1; Page 2394; 11750pp; English.

XX The invention relates to an isolated nucleic acid molecule (I) comprising  
 CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
 CC specification or its complement. (I) is useful for:  
 CC (a) assessing whether a patient is afflicted with prostate cancer;  
 CC (b) monitoring the progression of prostate cancer in a patient;  
 CC (c) assessing the efficacy of a test compound to inhibit prostate  
 CC cancer in a patient;  
 CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
 CC in a patient;  
 CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
 CC (f) assessing the prostate cell carcinogenic potential of a compound;  
 CC (g) determining whether prostate cancer has metastasized in a patient;  
 CC (h) assessing the aggressiveness or indolence of prostate cancer in a  
 CC patient;  
 CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.

SQ Sequence 192 BP; 38 A; 50 C; 70 G; 34 T; 0 other;

Query Match 69.2%; Score 16.6; DB 23; Length 192;  
 Best Local Similarity 82.6%; Pred. No. 98;  
 Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTCTCGTCGAGGCATCTAGT 24  
 DB 186 CTAGTCTCGTCGAGGCATCTCGT 164



Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker; pharmacogenomic marker; gene; ss.

Homo sapiens.

WO200160860-A2.

23-AUG-2001.

20-FEB-2001; 2001WO-US05171.

17-FEB-2000; 2000US-183319P.

16-MAR-2000; 2000US-189862P.

25-MAY-2000; 2000US-207454P.

09-JUN-2000; 2000US-211314P.

18-JUL-2000; 2000US-219007P.

13-DEC-2000; 2000US-255281P.

(MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.

Schlegel R, Endege WO, Monahan JE;

WPI; 2001-662795/76.

Novel isolated nucleic acid molecule associated with cancerous state of prostate cells and correlating with presence of prostate cancer, useful for detecting presence of prostate cancer, stage of prostate cancer -

Claim 1; Page 8790; 11750pp; English.

The invention relates to an isolated nucleic acid molecule (I) comprising a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the specification or its complement. (I) is useful for:

(a) assessing whether a patient is afflicted with prostate cancer;

(b) monitoring the progression of prostate cancer in a patient;

(c) assessing the efficacy of a test compound to inhibit prostate cancer in a patient;

(d) assessing the efficacy of a therapy for inhibiting prostate cancer in a patient;

(e) selecting a composition for inhibiting prostate cancer in a patient;

(f) assessing the prostate cell carcinogenic potential of a compound;

(g) determining whether prostate cancer has metastasized in a patient;

(h) assessing the aggressiveness or indolence of prostate cancer in a patient;

(I) is also useful as a pharmacodynamic or pharmacogenomic marker.

Sequence 450 BP; 93 A; 118 C; 143 G; 96 T; 0 other;

Query Match 69.2%; Score 16.6; DB 23; Length 450;

Best Local Similarity 82.6%; Pred. No. 1.1e+02;

Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTGTCTGCAGGATCTAGT 24

||||| ||||| ||||| ||

Db 228 CTAGTGTCTGCAGGATCTCGT 206

RESULT 14

ABV35433/C

ID ABV35433 standard; cDNA; 479 BP.

XX

AC ABV35433;

XX

16-SEP-2002 (first entry)

XX

Human prostate expression marker cDNA 35424.

XX

Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker; pharmacogenomic marker; gene; ss.

XX

Homo sapiens.

XX

WO200160860-A2.

PN

RESULT 12

AAC09061/C

ID AAC09061 standard; cDNA; 202 BP.

XX

AC AAC09061;

XX

06-OCT-2000 (first entry)

XX

Human secreted protein 5' EST, SEQ ID NO: 13136.

XX

Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation; gene therapy; chromosome mapping; ss.

XX

Homo sapiens.

XX

EP1033401-A2.

XX

06-SEP-2000.

XX

21-FEB-2000; 2000EP-0200610.

XX

26-FEB-1999; 99US-0122487.

XX

(GEST) GENSET.

XX

Dumas Milne Edwards J, Duclert A, Giordano J;

PI

WPI; 2000-500381/45.

DR

New nucleic acid that is a 5' expressed sequence tag (5' EST) for obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for diagnostic, forensic, gene therapy and chromosome mapping procedures -

XX

Claim 1; SEQ ID 13136; 71pp + CD-ROM; English.

XX

The present sequence is one of a large number of 5' ESTs derived from mRNAs encoding secreted proteins. No ORF has yet been conclusively identified within the present sequence. The 5' ESTs were prepared from total human RNAs or polyA+ RNAs derived from 30 different tissues. EST sequences usually correspond mainly to the 3' untranslated region (UTR) of the mRNA because they are often obtained from oligo-dT primed cDNA libraries. Such ESTs are not well suited for isolating cDNA sequences derived from the 5' ends of mRNAs and even in those cases where longer cDNA sequences have been obtained, the full 5' UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and chromosome mapping procedures. They are used to obtain upstream regulatory sequences and to design expression and secretion vectors.

XX

Sequence 202 BP; 40 A; 53 C; 74 G; 35 T; 0 other;

Query Match 69.2%; Score 16.6; DB 21; Length 202;

Best Local Similarity 82.6%; Pred. No. 99;

Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTGTCTGCAGGATCTAGT 24

||||| ||||| ||||| ||

Db 164 CTAGTGTCTGCAGGATCTCGT 142

RESULT 13

ABV44262/C

ID ABV44262 standard; cDNA; 450 BP.

XX

AC ABV44262;

XX

16-SEP-2002 (first entry)

XX

Human prostate expression marker cDNA 44253.

XX



GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:17 ; Search time 319.117 Seconds  
(without alignments)  
1218.024 Million cell updates/sec

Title: US-09-787-562-11

Perfect score: 24

Sequence: 1 tctagtctgtgcaggcatctagt 24

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

- EST:\*
- 1: em\_estba:\*
  - 2: em\_esthum:\*
  - 3: em\_estin:\*
  - 4: em\_estmu:\*
  - 5: em\_estov:\*
  - 6: em\_estpl:\*
  - 7: em\_estro:\*
  - 8: em\_htc:\*
  - 9: gb\_est1:\*
  - 10: gb\_est2:\*
  - 11: gb\_htc:\*
  - 12: gb\_est3:\*
  - 13: gb\_est4:\*
  - 14: gb\_est5:\*
  - 15: em\_estfun:\*
  - 16: em\_estom:\*
  - 17: gb\_gss:\*
  - 18: em\_gss\_hum:\*
  - 19: em\_gss\_inv:\*
  - 20: em\_gss\_pln:\*
  - 21: em\_gss\_vrt:\*
  - 22: em\_gss\_fun:\*
  - 23: em\_gss\_man:\*
  - 24: em\_gss\_mus:\*
  - 25: em\_gss\_other:\*
  - 26: em\_gss\_pro:\*
  - 27: em\_gss\_rod:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	20.4	85.0	529	BU011601	BU011601 QGJ16K24.
2	20.4	85.0	682	14 BQ843759	BQ843759 QGJ11I20.
3	18.2	75.8	567	17 BH206532	BH206532 Sml-46N18
4	17.8	74.2	297	17 AQ177064	AQ177064 HS-2216.B
5	17.8	74.2	503	17 BH729239	BH729239 BOMFY07TF
6	17.6	73.3	206	12 BF086292	BF086292 QV3-GN006

7	17.6	73.3	207	14 BQ366039	BQ366039 QV3-GN006
8	17.6	73.3	244	9 AA309771	AA309771 EST180630
9	17.6	73.3	300	17 AZ827575	AZ827575 2M0104005
10	17.6	73.3	520	12 BE930148	BE930148 QV3-GN006
11	17.6	73.3	582	9 AL705176	AL705176 DKF26868
12	17.6	73.3	607	17 AO882182	AO882182 HS-5358.B
13	17.6	73.3	750	12 BG830180	BG830180 602764757
14	17.6	73.3	812	9 AL521259	AL521259 AU521259
15	17.6	73.3	843	9 AU133085	AU133085 AU133085
16	17.6	73.3	875	9 AU125595	AU125595 AU125595
17	17.6	73.3	875	14 BQ215175	BQ215175 AGENCOURT
18	17.6	73.3	1016	13 BM543250	BM543250 AGENCOURT
19	17.6	73.3	1105	13 BM458582	BM458582 AGENCOURT
20	17.6	73.3	1194	12 BF780452	BF780452 602103972
21	17.2	71.7	109	10 AW994796	AW994796 RCI-BN003
22	17.2	71.7	283	13 BI007122	BI007122 QV3-RN006
23	17.2	71.7	389	17 AG019642	AG019642 Homo sapi
24	17.2	71.7	399	17 BH758471	BH758471 SALK 0199
25	17.2	71.7	417	14 BU024781	BU024781 QHFG606.Y
26	17.2	71.7	470	10 AW615392	AW615392 hg82d07.x
27	17.2	71.7	499	17 BH039133	BH039133 RPCI-24-2
28	17.2	71.7	544	17 AO678125	AO678125 HS-5528.B
29	17.2	71.7	561	17 AZ012741	AZ012741 RPCI-23-3
30	17.2	71.7	682	17 AO768930	AO768930 HS-3160.B
31	17.2	71.7	684	17 AG071473	AG071473 Pan trogl
32	17.2	71.7	712	12 BG431040	BG431040 602498853
33	17.2	71.7	832	17 AO897762	AO897762 HS-3153.A
34	17.2	71.7	879	14 BQ071181	BQ071181 AGENCOURT
35	17.2	71.7	891	12 BF672432	BF672432 ENTID73TF
36	17.2	71.7	1019	17 AZ688328	AZ688328 ENTID73TF
37	17.2	71.7	1113	13 BM043304	BM043304 603619567
38	17.2	71.7	1392	12 BF029841	BF029841 601556785
39	16.8	70.0	192	10 AM893929	AM893929 RCI-NN002
40	16.8	70.0	346	17 AZ774173	AZ774173 2M0003115
41	16.8	70.0	370	10 BE332565	BE332565 us24b04.x
42	16.8	70.0	433	10 BE335885	BE335885 us24b04.x
43	16.8	70.0	497	17 AQ298884	AQ298884 HS-3157.B
44	16.8	70.0	718	17 BH879219	BH879219 ht45a12.b
45	16.8	70.0	764	17 AQ870122	AQ870122 nbeb0036p

## ALIGNMENTS

RESULT 1	BU011601	529 bp	mRNA	linear	EST 22-AUG-2002
LOCUS	QGJ16K24.yg.ab1	QG_EFGHJ	lettuce	serriola	Lactuca sativa cDNA clone
DEFINITION	QGJ16K24, mRNA sequence.				
ACCESSION	BU011601				
VERSION	BU011601.1	GI:22445996			
KEYWORDS	EST				
SOURCE	Lactuca sativa.				
ORGANISM	Lactuca sativa				
REFERENCE	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;				
AUTHORS	Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Asteridae; euasterids II; Asterales; Asteraceae; Lactuceae; Lactuca				
	1 (bases 1 to 529)				
	Kozik, A., Michelmore, R.W., Knapp, S., Matvienko, M., Rieseberg, L., Lin, H., van Damme, M., Lavelle, D., Chevalier, P., Ziegler, J., Ellison, P., Kolkman, J., Slabaugh, M.S., Livingston, K., Zhou, Y., Lai, Z., Church, S., Jackson, L. and Bradford, K.				
TITLE	Lettuce and Sunflower ESTs from the Compositae Genome Project				
JOURNAL	http://compugenomics.ucdavis.edu/				
COMMENT	Unpublished (2002)				
	Contact: Alexander Kozik [R.W.Michelmore]				
	Department of Vegetable Crops, R.W.Michelmore Lab				
	University of California at Davis (UCD)				
	Asmundson Hall, UCD, Davis, CA 95616, USA				
	Tel: 1-(530)-742-1742				
	Fax: 1-(530)-752-9659				
	Email: akozik@atgc.org [michelmore@vegmail.ucdavis.edu]				

belongs to contig QG\_CA\_Contig2212, see <http://cgpdb.ucdavis.edu/> for details.  
 Plate: QG16 row: K column: 24.

#### FEATURES

Location/Qualifiers  
 1..529  
 /organism="Lactuca sativa"  
 /cultivar="L. serriola"  
 /db\_xref="taxon:4236"  
 /clone="QG16K24"  
 /clone\_lib="QG\_FRGHJ lettuce serriola"  
 /lab\_host="E.coli"  
 /note="Vector: pBRCDNASFIAB; The library was constructed from 10 different sources of RNA from a single genotype. Separate cDNAs were generated using primers that incorporated unique 5' and 3' tags to distinguish each source of RNA. cDNAs were then pooled, size-fractionated, directionally cloned into a custom medium-copy vector and transformations made with four size classes to minimize size bias. Details of each source of RNA and library construction can be obtained at <http://cgpdb.ucdavis.edu/>  
 TAG\_LIB=QG\_FRGHJ lettuce serriola  
 TAG\_TISSUE=flowers pre-fertilized  
 TAG\_SEQ=GCTTGACGGG"

BASE COUNT 182 a 83 c 115 g 149 t  
 ORIGIN

Query Match 85.0%; Score 20.4; DB 14; Length 529;

Best Local Similarity 95.5%; Pred. No. 64; Indels 0; Gaps 0;  
 Matches 21; Conservative 0; Mismatches 1;

QY 1 TCTAGTGTCTGCAGGCATCTA 22  
 ||||| ||||| ||||| ||||| |||||

Db 436 TCTAGTGTCTGCAGGCATCTA 457

#### RESULT 2

BQ843759

LOCUS

DEFINITION

VERSION

KEYWORDS

SOURCE

ORGANISM

Lactuca sativa

Lactuca sativa

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

Asteridae; euasterids II; Asterales; Asteraceae; Lactuceae;

Lactuca.

REFERENCE

AUTHORS

Kozik, A., Michelmore, R.W., Knapp, S., Matvienko, M., Rieseberg, L.,

Lin, H., van Damme, M., Lavelle, D., Chevalier, P., Ziegler, J., Ellison

P., Kolkman, J., Slabaugh, M.S., Livingston, K., Zhou, Y., Lai, Z.,

Church, S., Jackson, L. and Bradford, K.

Lettuce and Sunflower ESTs from the Compositae Genome Project

<http://compgenomics.ucdavis.edu/>

Unpublished (2002)

Contact: Alexander Kozik [R.W.Michelmore]

Department of Vegetable Crops, R.W.Michelmore Lab

University of California at Davis (UCD)

Asmundson Hall, UCD, Davis, CA 95616, USA

Tel: 1-(530)-742-1742

Fax: 1-(530)-752-9659

Email: akozik@ucdavis.edu [michelmore@vegmil.ucdavis.edu]

belongs to contig QG\_CA\_Contig2212, see <http://cgpdb.ucdavis.edu/> for details.

Plate: QG11 row: I column: 20.

Location/Qualifiers

1..682

/organism="Lactuca sativa"

/cultivar="Salinas"

/db\_xref="taxon:4236"

/clone="QG11I20"

#### FEATURES

source

/clone\_lib="QG\_ABCDI lettuce salinas"

/lab\_host="E.coli"

/note="Vector: pBRCDNASFIAB; The library was constructed from 10 different sources of RNA from a single genotype.

Separate cDNAs were generated using primers that

incorporated unique 5' and 3' tags to distinguish each

source of RNA. cDNAs were then pooled, size-fractionated,

directionally cloned into a custom medium-copy vector and

transformations made with four size classes to minimize

size bias. Details of each source of RNA and library

construction can be obtained at <http://cgpdb.ucdavis.edu/>  
 TAG\_SEQ=Not found"

BASE COUNT 210 a 119 c 157 g 196 t

ORIGIN

Query Match 85.0%; Score 20.4; DB 14; Length 682;

Best Local Similarity 95.5%; Pred. No. 71;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGCAGGCATCTA 22

||||| ||||| ||||| ||||| |||||

Db 585 TCTAGTGTCTGCAGGCATCTA 606

#### RESULT 3

BH206532/c

LOCUS

DEFINITION

SEQUENCE

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Schistosoma mansoni

Schistosoma mansoni

Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;

Strigeidida; Schistosomatoidea; Schistosomatidae; Schistosoma.

REFERENCE

AUTHORS

Shetty, J., Simpson, A., Malek, J., Koo, H., Loverde, P.T. and El-Sayed

, N.M.

Use of end sequences from Schistosoma mansoni (Puerto Rico strain)

Sml BAC library for gene discovery and map construction

Unpublished (2001)

Other\_GSSs: Sml-46N18.TF

Contact: Najib M. El-Sayed

Department of Eukaryotic Genomics

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 0200

Fax: 301 838 0208

Email: nelsayed@tigr.org

10.edu).

Seq primer: M13 Rev

Class: BAC ends.

Location/Qualifiers

1..567

/organism="Schistosoma mansoni"

/strain="Puerto Rico"

/db\_xref="taxon:6183"

/clone="Sml-46N18"

/clone\_lib="Sml"

/note="Vector: pBelOBA11; Site 1: Hin dIII; Constructed

in the laboratory of Dr. Denis Le Paslier at the Fondation

Jean Dausset, CEPH, Paris, France. Briefly, Schistosoma

mansoni agarose embedded DNA was partially digested with

Hin dIII. High molecular weight fragments were ligated in

pBelOBA11 digested with Hin dIII. The average insert size

is 100 kb. Total clone coverage: approx. 7.95 x the

haploid genome. Further information can be found in Le

Paslier et al. (2000) Construction and characterization of

a Schistosoma mansoni bacterial artificial chromosome

library. Genomics 65: 87-94."

BASE COUNT 197 a 108 c 114 g 148 t

ORIGIN

Brazil  
Readalicia Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project This entry can be seen in the following URL  
(<http://www.ludwig.org.br/scripts/gethtml2.pl?tl=st3-QV3-GN0065-120>)  
900-323-50&st3=2000-09-12&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 6

High quality sequence stop: 206.

## FEATURES

source Location/Qualifiers

1..206  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_lib="GN0065"  
/dev\_stage="Adult"  
/note="Organ: Placenta\_normal; Vector: puc18; Site\_1: SmaI  
; Site\_2: SmaI; A mini-library was made by cloning  
products derived from ORESTES PCR (U.S. Letters Patent  
application No. 196,716 - Ludwig Institute for Cancer  
Research) profiles into the pUC 18 vector. Reverse  
transcription of tissue mRNA and cDNA amplification were  
performed under low stringency conditions."

63 a 37 c 51 g 55 t

## BASE COUNT

ORIGIN

Query Match 73.3%; Score 17.6; DB 12; Length 206;

Best Local Similarity 83.3%; Pred. No. 8e+02; Indels 0; Gaps 0;

Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCACTAGT 24

||||| ||||| ||||| |||||

Db 94 TCTGGAGTCGTGGAGCACTAGT 117

## RESULT 7

BQ366039

LOCUS

DEFINITION

QV3-GN0061-270900-323-g02 GN0061 Homo sapiens cDNA, mRNA sequence.

ACCESSION

BQ366039

VERSION

BQ366039.1 GI:21041551

KEYWORDS

EST.

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 207)

Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R.,

Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,

Goldman, G.H., Carvalho, A.F., Matsukuma, A., Baia, G.S., Simpson, D.H.,

Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare

, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and

Simpson, A.J.

Shotgun sequencing of the human transcriptome with ORF expressed

sequence tags

Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

20202663

Contact: Simpson A.J.G.

Laboratory of Cancer Genetics

Ludwig Institute for Cancer Research

Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,

Brazil

Tel: +55-11-2704922

Fax: +55-11-2707001

Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome

project. This entry can be seen in the following URL

(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=QV3&t2=QV3-GN0061-

270900-323-g02&t3=2000-09-27&t4=1)

Seq primer: puc 18 forward

High quality sequence stop: 6

High quality sequence stop: 207.

Location/Qualifiers

1..207

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone\_lib="GN0061"

/dev\_stage="Adult"

/note="Organ: placenta\_normal; Vector: puc18; Site\_1: SmaI

; Site\_2: SmaI; A mini-library was made by cloning

products derived from ORESTES PCR (U.S. Letters Patent

application No. 196,716 - Ludwig Institute for Cancer

## BASE COUNT

ORIGIN

Query Match 73.3%; Score 17.6; DB 14; Length 207;

Best Local Similarity 83.3%; Pred. No. 8e+02; Indels 0; Gaps 0;

Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCACTAGT 24

||||| ||||| ||||| |||||

Db 95 TCTGGAGTCGTGGAGCACTAGT 118

## RESULT 8

AA309771

LOCUS

DEFINITION

EST180630 Jurkat T-cells V Homo sapiens cDNA 5' end similar to

hypothetical protein KIAA0225, mRNA sequence.

AA309771

ACCESSION

AA309771.1 GI:1962100

VERSION

EST.

KEYWORDS

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 244)

Adams, M.D., Kerlavage, A.R., Fleischmann, R.D., Fuldner, R.A., Bult

, C.J., Lee, N.H., Kirkness, E.F., Weinstock, K.G., Gocayne, J.D., White

, O., Sutton, G., Blake, J.A., Brandon, R.C., Man-Wai, C., Clayton, R.A.,

Cline, T.R., Cotton, M.D., Earle-Hughes, J., Fine, L.D., Fitzgerald

, L.M., Fitzhugh, W.M., Fritchman, J.L., Geoghagen, N.S., Glodek, A.,

Gnehm, C.L., Hanna, M.C., Hedblom, E., Hinkle, P.S. Jr., Kelley, J.M.,

Kelley, J.C., Liu, L.-I., Marmaros, S.M., Merrick, J.M.,

Moreno-Palancas, R.F., McDonald, L.A., Nguyen, D.T., Pelligrino, S.M.,

Phillips, C.A., Ryder, S.E., Scott, J.L., Saudek, D.M., Shirley, R.,

Small, K.V., Spriggs, F.A., Utterback, T.R., Weidman, J.F., Li, Y.,

Bednarik, D.P., Cao, L., Cepeda, M.A., Coleman, T.A., Collins, E.J.,

Dimke, D., Feng, D.-F., Ferrie, A., Fischer, C., Hastings, G.A., He, W.W.,

Hu, J.S., Greene, J.M., Gruber, J., Hudson, P., Kim, A.K., Kozak, D.L.,

Kunsch, C., Greene, J.M., Gruber, J., Hudson, P., Kim, A.K., Kozak, D.L.,

Wei, Y.F., Wang, J., Xu, C., Yu, G.L., Ruben, S.M., Dillion, P.J., Fannon

, M.R., Rosen, C.A., Haseltine, W.A., Fields, C., Fraser, C.M. and

Venter, J.C.

Initial assessment of human gene diversity and expression patterns

based upon 83 million nucleotides of cDNA sequence

Nature 377 (6547 Suppl), 3-174 (1995)

96026280

Other\_ESTs: THCL89719

Contact: Kerlavage, AR

Bioinformatics

The Institute for Genomic Research

9712 Medical Center Drive, Rockville, MD 20850 USA

Tel: 3018699056

Fax: 3018699423

Email: arkerlav@tigr.org

For clone availability, additional sequence and expression

information related to this EST, please check the tigr Human Gene

Index (http://www.tigr.org/tdb/hgi/hgi.html)

Seq primer: M3 Reverse.

Location/Qualifiers

1..244

/organism="Homo sapiens"

/db\_xref="ATCC (inhost):155863"

/db\_xref="taxon:9606"

/clone\_lib="Jurkat T-cells V"

/cell\_type="T-lymphocyte"

/note="Vector: pBluescript SK-; Site\_1: EcoRI; Site\_2:

XhoI"

73 a 46 c 59 g 63 t 3 others

BASE COUNT

ORIGIN

Research) profiles into the pUC 18 vector. Reverse  
transcription of tissue mRNA and cDNA amplification were  
performed under low stringency conditions."

64 a 37 c 51 g 55 t

Query Match 73.3%; Score 17.6; DB 14; Length 207;

Best Local Similarity 83.3%; Pred. No. 8e+02; Indels 0; Gaps 0;

Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCACTAGT 24

||||| ||||| ||||| |||||

Db 95 TCTGGAGTCGTGGAGCACTAGT 118

BASE COUNT  
ORIGIN

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LOCUS      AL705176      582 bp      mRNA      linear      EST 22-MAR-2002
DEFINITION DKFZp686N1034_r1 686 (synonym: hlcc3) Homo sapiens cDNA clone
ACCESSION  AL705176
VERSION     AL705176.1 GI:19688531
KEYWORDS   EST.
SOURCE      human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 582)
AUTHORS     Ottenwaelder,B., Obermaier,B., Mewes,H.W., Mewes,H.W., Weil,B. and
            Wiemann,S.
TITLE       EST (Ottenwaelder,B., Obermaier,B., Mewes,H.W., Weil,B. and Wiemann
            S.)
JOURNAL     Unpublished (2001)
COMMENT     Contact: Ottenwaelder B
            MIPS
            Am Klopferstritz 18a D-82152 Martinsried, Germany
            This is the 5' sequence of the clone insert
            Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
            Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de;
            sequenced by Medigenomix (Martinsried/Germany) within the cDNA
            sequencing consortium of the German Genome Project. No sl sequence
            available.
            This clone (DKFZp686N1034) is available at the RZPD in Berlin.
            Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
            Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.
FEATURES   source
            Location/Qualifiers
                1..582
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="DKFZp686N1034"
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                /tissue_type="human skeletal muscle"
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BASE COUNT  168 a 123 c 137 g 154 t
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Query Match 73.3%; Score 17.6; DB 9; Length 582;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TCTAGTCTGTCGAGGCATCTAGT 24
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Db 491 TCTGGAGTCTGTCGAGGCAACTAGT 514

RESULT 12
LOCUS      AQ882182      607 bp      DNA      linear      GSS 09-NOV-1999
DEFINITION HS_5358_B2_B06_T7A RPCI-11 Human Male BAC Library Homo sapiens
            genomic clone Plate=9126 Col=12 Row=D, DNA sequence.
ACCESSION  AQ882182
VERSION     AQ882182.1 GI:6313649
KEYWORDS   GSS.
SOURCE      human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 607)
AUTHORS     Mahairas,G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
            Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
            Hood,L.
TITLE       Sequence-tagged connectors: A sequence approach to mapping and
            scanning the human genome
JOURNAL     Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE     99380589
COMMENT     Contact: Mahairas GG, Wallace JC, Hood L
            High Throughput Sequencing Center

```

```

University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 9126 row: D column: 12
Seq primer: r7
Class: BAC ends
High quality sequence stop: 607.
FEATURES   source
            Location/Qualifiers
                1..607
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="Plate=9126 Col=12 Row=D"
                /clone_lib="RPCI-11 Human Male BAC Library"
                /sex="male"
                /note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
                Male blood DNA was isolated from one randomly chosen donor
                and partially digested with a combination of EcoRI and
                EcoRI Methylase. Size selected DNA was cloned into the
                pBACe3.6 vector at EcoRI sites"
BASE COUNT  170 a 116 c 109 g 204 t
ORIGIN
Query Match 73.3%; Score 17.6; DB 17; Length 607;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TCTAGTCTGTCGAGGCATCTAGT 24
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Db 231 TCTAGTCTGTCGATGATCTAGT 254

RESULT 13
LOCUS      BG830180      750 bp      mRNA      linear      EST 22-MAY-2001
DEFINITION 602764757F1 NIH_MGC_42 Homo sapiens cDNA clone IMAGE:4906671 5',
            mRNA sequence.
ACCESSION  BG830180
VERSION     BG830180.1 GI:14177767
KEYWORDS   EST.
SOURCE      human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 750)
AUTHORS     NIH-MGC http://mgc.nci.nih.gov/.
            National Institutes of Health, Mammalian Gene Collection (MGC)
            Unpublished (1999)
JOURNAL
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgabs-r@mail.nih.gov
            Tissue Procurement: ATCC
            cDNA Library Preparation: Ling Hong/Rubin Laboratory
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLCMI810 row: b column: 16
            High quality sequence stop: 723.
FEATURES   source
            Location/Qualifiers
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                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="IMAGE:4906671"
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/lab\_host="DH10B (phage-resistant)"  
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 Site.2: EcoRI; cDNA made by oligo-dT priming.  
 Directionally cloned into EcoRI/XhoI sites using the  
 following 5' adaptor: GGCACGAG(G). Size-selected >500bp  
 for average insert size 1.8Kb. Library constructed by Ling  
 Hong in the laboratory of Gerald M. Rubin (University of  
 California, Berkeley) using ZAP-cDNA synthesis kit  
 (Stratagene) and Superscript II RT (Life Technologies).  
 Note: this is a NIH\_MGC Library. |"  
 BASE COUNT 204 a 159 c 182 g 205 t  
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Query Match 73.3%; Score 17.6; DB 12; Length 750;  
 Best Local Similarity 83.3%; Pred. No. 1.3e+03;  
 Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
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 Db 279 TCTGGAGTCGTGGAGGCAACTAGT 302

RESULT 14  
 AL521259 812 bp mRNA linear EST 13-FEB-2001  
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 DEFINITION prime, mRNA sequence.  
 ACCESSION AL521259  
 VERSION AL521259.1 GI:12784752  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 812)  
 AUTHORS Li, W.B., Gruber, C., Jessee, J. and Polayes, D.  
 TITLE Full-length cDNA libraries and normalization  
 JOURNAL Unpublished (2001)  
 COMMENT Contact: Genoscope  
 Genoscope - Centre National de Sequencage  
 BP 191 91006 EVRY cedex - France  
 Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.  
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 /clone\_lib="LTI\_NFL004\_NBC2"  
 /sex="male"  
 /tissue\_type="neuroblastoma cells"  
 /lab\_host="DH10B"  
 /note="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA  
 was primed with a NotI-oligo(dT) primer. Five prime end  
 enriched, double-stranded cDNA was digested with Not I and  
 cloned into the Not I and Eco RV sites of the pCMVSPORT 6  
 vector. Library was normalized. Library was constructed  
 by Life Technologies. Contact : Feng Liang Life  
 Technologies, a division of Invitrogen 9800 Medical Center  
 Drive Rockville, Maryland 20850, USA Fax : (1) 301 610  
 8371 Email : fliang@lifetech.com URL :  
 http://fulllength.invitrogen.com"  
 BASE COUNT 220 a 174 c 195 g 221 t 2 others  
 ORIGIN

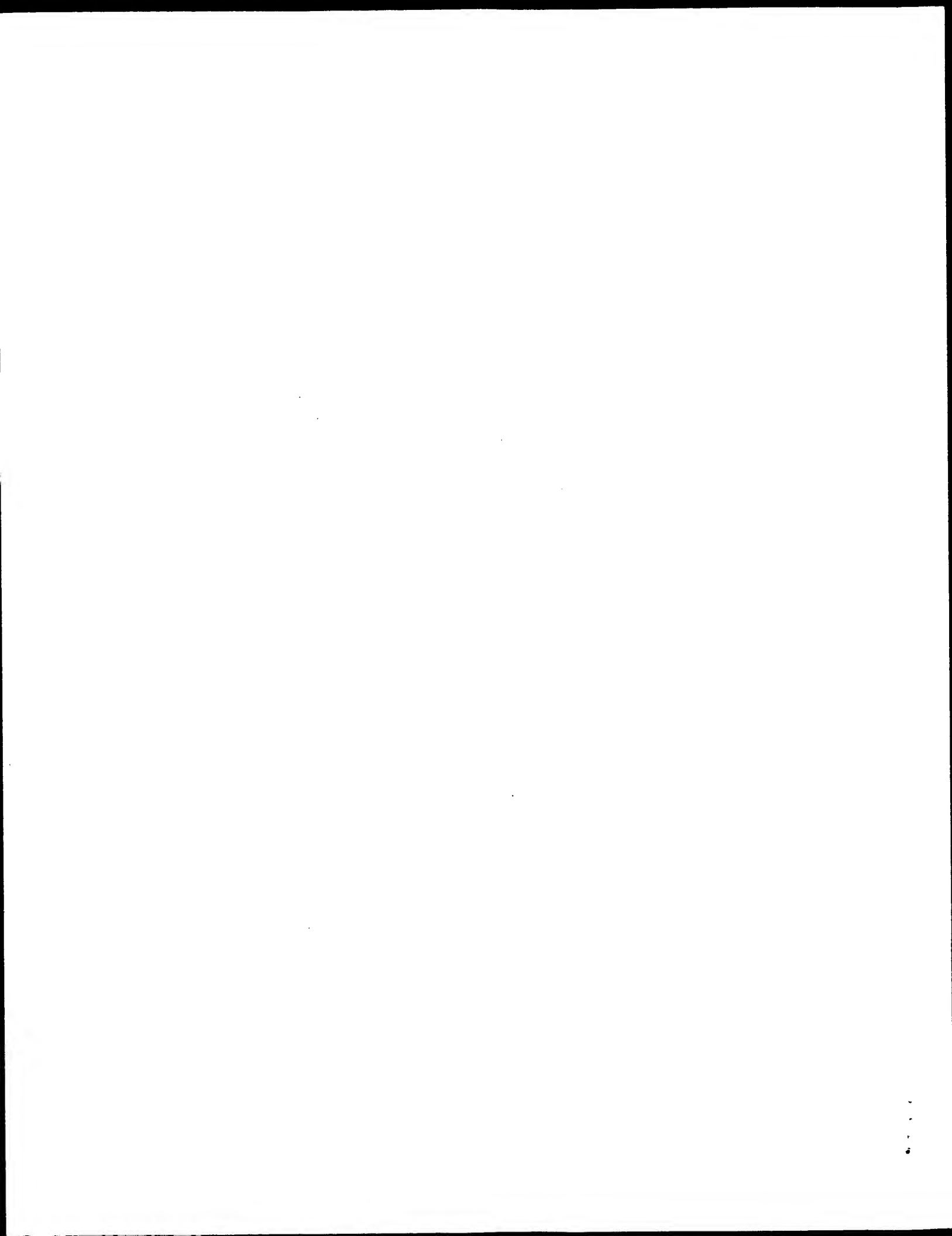
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 Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 1 TCTAGTGTCTGTCGAGGCATCTAGT 24  
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RESULT 15  
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 DEFINITION sequence.  
 ACCESSION AUI33085  
 VERSION AUI33085.1 GI:10993624  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 843)  
 AUTHORS Ota, T., Sugiyama, T., Ishii, S., Suzuki, Y., Saito, K., Yamamoto, J.,  
 Nishikawa, T., Nakamura, Y., Nagai, T., Sugano, S., Masuho, Y. and  
 Isogai, T.  
 TITLE HRI human cDNA project (Ota, T., Sugiyama, T., Ishii, S., Suzuki, Y.,  
 Saito, K., Yamamoto, J., Nishikawa, T., Nakamura, Y., Nagai, T., Sugano  
 S., Masuho, Y., Isogai, T.)  
 JOURNAL Unpublished (2000)  
 COMMENT Contact: Takao Isogai  
 Genomics Laboratory  
 Helix Research Institute  
 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan  
 Tel: 81-438-52-3975  
 Fax: 81-438-52-3986  
 Email: genomics@hri.co.jp  
 HRI human cDNA project; 5' - & 3'-end one pass sequencing: Helix  
 Research Institute; cDNA library construction: Department of  
 Virology, Institute of Medical Science, University of Tokyo, and  
 Helix Research Institute.  
 Location/Qualifiers  
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 /db\_xref="taxon:9606"  
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 ORIGIN

Query Match 73.3%; Score 17.6; DB 9; Length 843;  
 Best Local Similarity 83.3%; Pred. No. 1.4e+03;  
 Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 1 TCTAGTGTCTGTCGAGGCATCTAGT 24  
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 Db 188 TCTGGAGTCGTGGAGGCAACTAGT 211

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 Job time : 325.117 secs



GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:53:16 ; Search time 8.10095 Seconds  
(without alignments)  
908.566 Million cell updates/sec

Title: US-09-787-562-11

Perfect score: 24

Sequence: 1 tctagtctgtcgcaggtatctagt 24

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents.NA.\*

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- 6: /cgn2\_6/ptodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

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2	16.6	69.2	3807	2	US-09-003-289-1
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4	16.6	69.2	7559	2	US-08-250-848-2
5	16.6	66.7	17656	4	US-09-433-579-3
6	15.4	64.2	2057	3	US-09-008-303-1
7	15.4	64.2	246240	2	US-08-724-394A-20
8	15.4	64.2	246240	2	US-08-724-394A-21
9	15.4	64.2	246240	2	US-08-724-394A-22
10	15.2	63.3	2219	1	US-08-290-979A-7
11	15	62.5	594	4	US-09-328-111-155
12	15	62.5	1128	1	US-08-448-744-6
13	15	62.5	17056	4	US-09-245-041-3
14	14.8	61.7	822	1	US-07-644-372-1
15	14.8	61.7	3448	1	US-08-296-014A-3
16	14.8	61.7	3448	1	US-08-596-405-3
17	14.8	61.7	3448	2	US-08-877-620-3
18	14.8	61.7	4173	4	US-08-981-729-9
19	14.8	61.7	4173	4	US-08-981-446B-2
20	14.8	61.7	4182	1	US-08-296-014A-1
21	14.8	61.7	4182	2	US-08-596-405-1
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23	14.6	60.8	302	3	US-08-746-111-36
24	14.6	60.8	1251	4	US-09-072-596-319
25	14.6	60.8	3097	2	US-08-599-455B-1
26	14.6	60.8	3097	4	US-09-069-781B-1
27	14.6	60.8	3097	4	US-09-137-132-1

28	14.6	60.8	3097	4	US-08-864-564A-1	Sequence 1, Appli
29	14.6	60.8	3097	4	US-09-094-410-1	Sequence 1, Appli
30	14.6	60.8	3854	2	US-08-599-455B-42	Sequence 42, Appl
31	14.6	60.8	3854	4	US-09-069-781B-42	Sequence 42, Appl
32	14.6	60.8	3854	4	US-09-137-132-42	Sequence 42, Appl
33	14.6	60.8	3854	4	US-08-864-564A-42	Sequence 42, Appl
34	14.6	60.8	3854	4	US-09-094-410-42	Sequence 42, Appl
35	14.6	60.8	4264	2	US-08-784-649A-1	Sequence 1, Appli
36	14.6	60.8	4264	2	US-08-784-649A-5	Sequence 5, Appli
37	14.6	60.8	4646	1	US-08-181-471-2	Sequence 2, Appli
38	14.6	60.8	4669	2	US-08-583-276-18	Sequence 18, Appl
39	14.6	60.8	4669	2	US-08-752-447-1	Sequence 1, Appli
40	14.6	60.8	4669	4	US-09-316-167-1	Sequence 1, Appli
41	14.6	60.8	4669	6	5206352-3	Patent No. 5206352
42	14.6	60.8	4773	3	US-08-884-324-9	Sequence 9, Appli
43	14.6	60.8	6505	2	US-08-793-610-5	Sequence 5, Appli
44	14.6	60.8	6737	4	US-09-453-702B-76	Sequence 76, Appl
45	14.6	60.8	9318	2	US-08-793-610-6	Sequence 6, Appli

## ALIGNMENTS

RESULT 1  
US-08-357-598-1  
; Sequence 1, Application US/08357598  
; Patent No. 5705625  
; GENERAL INFORMATION:  
; APPLICANT: Civin, Curt I.  
; APPLICANT: Small, Donald  
; TITLE OF INVENTION: NOVEL PROTEIN TYROSINE KINASE, JAK3  
; NUMBER OF SEQUENCES: 12  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Fish & Richardson P.C.  
; STREET: 4225 Executive Square, Suite 1400  
; CITY: La Jolla  
; STATE: CA  
; COUNTRY: USA  
; ZIP: 92037  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/357,598  
; FILING DATE: 15-DEC-1994  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Halle, Lisa A.  
; REGISTRATION NUMBER: 38,347  
; REFERENCE/DOCKET NUMBER: 07265/033001  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 619/678-5070  
; TELEFAX: 619/678-5099  
; INFORMATION FOR SEQ ID NO: 1:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 3807 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA  
US-08-357-598-1

Query Match 69.2%; Score 16.6; DB 1; Length 3807;  
Best Local Similarity 82.6%; Pred. No. 26;  
Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
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Db 1896 CAAGTCTCTGCGAGGCATCTCGT 1918

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RESULT 2
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; Sequence 1, Application US/09003289
; Patent No. 5916792
; GENERAL INFORMATION:
; APPLICANT: Civin, Curt I.
; APPLICANT: Small, Donald
; TITLE OF INVENTION: NOVEL PROTEIN TYROSINE KINASE, JAK3
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: USA
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/003,289
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/357,598
; FILING DATE: 15-DEC-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Haile, Lisa A.
; REGISTRATION NUMBER: 38,347
; REFERENCE/DOCKET NUMBER: 07265/033001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619/678-5099
; TELEFAX: 619/678-5099
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3807 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
US-09-003-289-1
Query Match 69.2%; Score 16.6; DB 2; Length 3807;
Best Local Similarity 82.6%; Pred. No. 26;
Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTGTCTGTCGAGGCATCTAGT 24
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Db 1896 CAAAGTGTCTGTCACCGCATCTCGT 1918

RESULT 3
PCT-US95-16435-1
; Sequence 1, Application PC/TUS9516435
; GENERAL INFORMATION:
; APPLICANT: The Johns Hopkins University School of Medicine
; TITLE OF INVENTION: NOVEL PROTEIN TYROSINE KINASE, JAK3
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: USA
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
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APPLICATION NUMBER: PCT/US95/16435
FILING DATE: 15-DEC-1995
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07265/033WO1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 3807 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
PCT-US95-16435-1
Query Match 69.2%; Score 16.6; DB 5; Length 3807;
Best Local Similarity 82.6%; Pred. No. 26;
Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTGTCTGTCGAGGCATCTAGT 24
| | | | | | | | | | | | | | | |
Db 1896 CAAAGTGTCTGTCACCGCATCTCGT 1918

RESULT 4
US-08-250-848-2
; Sequence 2, Application US/08250848
; Patent No. 5856177
; GENERAL INFORMATION:
; APPLICANT: Hudspeth, Richard L.
; TITLE OF INVENTION: PROMOTERS DERIVED FROM THE MAIZE
; TITLE OF INVENTION: PHOSPHOENOLPYRUVATE CARBOXYLASE GENE INVOLVED IN C4
; TITLE OF INVENTION: PHOTOSYNTHESIS
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Christie, Parker & Hale
; STREET: P.O. Box 7068
; CITY: Pasadena
; STATE: CA
; COUNTRY: USA
; ZIP: 91009-7068
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/250,848
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Sharp Esq., Janice A.
; REGISTRATION NUMBER: 34,051
; REFERENCE/DOCKET NUMBER: P114:25992/JAS
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (818) 795-5843
; TELEFAX: (818) 577-1769
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7559 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Zea mays
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; LOCATION: 4492..4596
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 4723..4812
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 4945..5100
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 5198..6196
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 6294..6680
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 6789..7079
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; LOCATION: 2153..2332
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; LOCATION: 2437..2832
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; LOCATION: 3901..4120
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; NAME/KEY: intron
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; FEATURE:
; NAME/KEY: intron
; LOCATION: 4946..5099
; FEATURE:
; NAME/KEY: intron
; LOCATION: 5199..6195
; FEATURE:
; NAME/KEY: intron
; LOCATION: 6295..6679
; FEATURE:
; NAME/KEY: intron
; LOCATION: 6790..7078
; FEATURE:
; NAME/KEY: polyA_site
; LOCATION: 7314..7319
; FEATURE:
; NAME/KEY: prim_transcript
; LOCATION: 2072
; FEATURE:
; NAME/KEY: promoter
; LOCATION: 1..2155
; FEATURE:
; NAME/KEY: TATA_signal
; LOCATION: 2042..2049
; FEATURE:
; NAME/KEY: mRNA
; LOCATION: 2153..7079
; US-08-250-848-2
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Query Match 69.2%; Score 16.6; DB 2; Length 7559;
Best Local Similarity 82.6%; Pred. No. 28;
Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
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QY 2 CTAGTGTCTGTCAGGCATCTAGT 24
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Db 5176 CTACTGTCTGTCATGTCAGT 5198
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RESULT 5
US-09-433-579-3/c
; Sequence 3, Application US/09433579
; Patent No. 6444877
; GENERAL INFORMATION:
; APPLICANT: Rottmann, William H.
; TITLE OF INVENTION: LSAG Gene
; FILE REFERENCE: LSAG Gene
; CURRENT APPLICATION NUMBER: US/09/433,579
; CURRENT FILING DATE: 1999-11-04
; NUMBER OF SEQ ID NOS: 36
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
; LENGTH: 17656
; TYPE: DNA
; ORGANISM: Liquidambar styraciflua
; US-09-433-579-3
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Query Match 66.7%; Score 16; DB 4; Length 17656;
Best Local Similarity 79.2%; Pred. No. 61;
Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
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QY 1 TCTAGTGTCTGTCAGGCATCTAGT 24
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Db 15206 TCTAGTGGGTGCAAGCTGCTAGT 15183
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RESULT 6
US-09-008-303-1/c
; Sequence 1, Application US/09008303
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; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/724,394A  
; FILING DATE: 01-OCT-1996  
; CLASSIFICATION: 536  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Fitts, Renee A.  
; REGISTRATION NUMBER: 35,136  
; REFERENCE/DOCKET NUMBER: 017957-000100  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 415-576-0200  
; TELEFAX: 415-576-0300  
; INFORMATION FOR SEQ ID NO: 21:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 246240 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: not relevant  
; TOPOLOGY: not relevant  
; MOLECULE TYPE: cdna  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: 1..246240  
; OTHER INFORMATION: /note= "HLA-H.CONTIG"  
US-08-724-394A-21

Query Match 64.2%; Score 15.4; DB 2; Length 246240;  
Best Local Similarity 94.1%; Pred. No. 1.6e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 5 GTGTCGTGCGAGCATCT 21  
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Db 142670 GTGTCGTGCGAGCATCT 142686

RESULT 9  
US-08-724-394A-22  
; Sequence 22, Application US/08724394A  
; Patent No. 5872237  
; GENERAL INFORMATION:  
; APPLICANT: Feder, John N.  
; APPLICANT: Kronmal, Gregory S.  
; APPLICANT: Lauer, Peter M.  
; APPLICANT: Ruddy, David A.  
; APPLICANT: Thomas, Winston  
; APPLICANT: Tsuchihashi, Zenta  
; APPLICANT: Wolff, Roger K.  
; TITLE OF INVENTION: Megabase Transcript Map: No. 5872237el  
; NUMBER OF SEQUENCES: 31  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: TOWNSEND and TOWNSEND and CREW LLP  
; STREET: Two Embarcadero Center, 8th Floor  
; CITY: San Francisco  
; STATE: CA  
; COUNTRY: USA  
; ZIP: 94111-3834  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/724,394A  
; FILING DATE: 01-OCT-1996  
; CLASSIFICATION: 536  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Fitts, Renee A.  
; REGISTRATION NUMBER: 35,136  
; REFERENCE/DOCKET NUMBER: 017957-000100  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 415-576-0200  
; TELEFAX: 415-576-0300  
; INFORMATION FOR SEQ ID NO: 22:

; SEQUENCE CHARACTERISTICS:  
; LENGTH: 246240 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: not relevant  
; TOPOLOGY: not relevant  
; MOLECULE TYPE: cdna  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: 1..246240  
; OTHER INFORMATION: /note= "HLA-H.CONTIG"  
US-08-724-394A-22

Query Match 64.2%; Score 15.4; DB 2; Length 246240;  
Best Local Similarity 94.1%; Pred. No. 1.6e+02;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 5 GTGTCGTGCGAGCATCT 21  
|||||  
Db 142670 GTGTCGTGCGAGCATCT 142686

RESULT 10  
US-08-290-979A-7/c  
; Sequence 7, Application US/08290979A  
; Patent No. 5610046  
; GENERAL INFORMATION:  
; APPLICANT: VAN OOIJEN, Albert J.H.  
; APPLICANT: DE GRAAFF, Leendert H.  
; APPLICANT: VAN DEN BROECK, Henriette C.  
; APPLICANT: VISSER, Jacob  
; TITLE OF INVENTION: Cloning and Expression of Xylanase B  
; NUMBER OF SEQUENCES: 8  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: MORRISON & FOERSTER  
; STREET: 2000 Pennsylvania Ave. N.W.  
; CITY: Washington  
; STATE: D.C.  
; COUNTRY: USA  
; ZIP: 20006-1812  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/290,979A  
; FILING DATE: 22-SEP-1994  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: KATE H. MURASHIGE  
; REGISTRATION NUMBER: 29,959  
; REFERENCE/DOCKET NUMBER: 4615-0045.00  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (202) 887-1500  
; TELEFAX: (202) 887-0763  
; TELEX: 90-4030 MRSN FOERS WSH  
; INFORMATION FOR SEQ ID NO: 7:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2219 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; HYPOPHETICAL: NO  
; ORIGINAL SOURCE:  
; ORGANISM: Aspergillus tubigenis  
; STRAIN: DSI6813  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: join(902..1180, 1248..1643)  
US-08-290-979A-7

Query Match 63.3%; Score 15.2; DB 1; Length 2219;

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Best Local Similarity 85.0%; Pred. No. 1.2e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCATCTC 20
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Db 624 TCTAGGTCTCTGCAGACATC 605

RESULT 11
US-09-328-111-155
; Sequence 155, Application US/09328111
; Patent No. 6262333
; GENERAL INFORMATION:
; APPLICANT: Endege, Wilson O.
; APPLICANT: Steinmann, Kathleen E.
; APPLICANT: Astle, Jon H.
; APPLICANT: Burgess, Christopher C.
; APPLICANT: Bushnell, Steven E.
; APPLICANT: Carroll III, Eddie
; APPLICANT: Catino, Theodore J.
; APPLICANT: Derti, Adnan
; APPLICANT: Ford, Donna M.
; APPLICANT: Lewis, Marcia E.
; APPLICANT: Monahan, John E.
; APPLICANT: Schlegel, Robert
; TITLE OF INVENTION: NOVEL HUMAN GENES AND GENE EXPRESSION
; TITLE OF INVENTION: PRODUCTS
; FILE REFERENCE: CCD-257 (US)
; CURRENT APPLICATION NUMBER: US/09/328,111
; CURRENT FILING DATE: 1999-06-08
; EARLIER APPLICATION NUMBER: US 60/088,801
; EARLIER FILING DATE: 1998-06-10
; NUMBER OF SEQ ID NOS: 850
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 155
; LENGTH: 594
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(594)
; OTHER INFORMATION: n = A,T,C or G
US-09-328-111-155

Query Match 62.5%; Score 15; DB 4; Length 594;
Best Local Similarity 75.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCATCTAGT 24
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Db 412 TCAAGTGTGTGCTTCATCTCCT 435

RESULT 12
US-08-448-744-6/C
; Sequence 6, Application US/08448744
; Patent No. 5616484
; GENERAL INFORMATION:
; APPLICANT: XU, Shuang-yong
; TITLE OF INVENTION: Cloning And Expression of The ApaLI
; TITLE OF INVENTION: Restriction Endonuclease
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: New England Biolabs, Inc.
; STREET: 32 Tozer Road
; CITY: Beverly
; STATE: Massachusetts
; COUNTRY: US
; ZIP: 01915
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS

Best Local Similarity 85.0%; Pred. No. 1.2e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCATC 20
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Db 624 TCTAGGTCTCTGCAGACATC 605

SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/448,744
FILING DATE: 24-May-1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Williams, Gregory D.
REGISTRATION NUMBER: 30901
REFERENCE/DOCKET NUMBER: NEB-114
TELECOMMUNICATION INFORMATION:
TELEPHONE: (508) 927-5054
TELEFAX: (508) 927-1705
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 1128 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
US-08-448-744-6

Query Match 62.5%; Score 15; DB 1; Length 1128;
Best Local Similarity 78.3%; Pred. No. 1.4e+02;
Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCATCTAG 23
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Db 302 TCCAGTGAAGTGCCTGCATCAAG 280

RESULT 13
US-09-245-041-3
; Sequence 3, Application US/09245041
; Patent No. 6274339
; GENERAL INFORMATION:
; APPLICANT: Moore, K.
; APPLICANT: Nagle, D.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DIAGNOSIS AND TREATMENT
; TITLE OF INVENTION: OF BODY WEIGHT DISORDERS INCLUDING OBESITY
; FILE REFERENCE: 7853-136
; CURRENT APPLICATION NUMBER: US/09/245,041
; CURRENT FILING DATE: 1999-02-05
; EARLIER APPLICATION NUMBER: 60/093,630
; EARLIER FILING DATE: 1998-07-21
; EARLIER APPLICATION NUMBER: 60/104,978
; EARLIER FILING DATE: 1998-10-20
; NUMBER OF SEQ ID NOS: 131
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 3
; LENGTH: 17056
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-245-041-3

Query Match 62.5%; Score 15; DB 4; Length 17056;
Best Local Similarity 78.3%; Pred. No. 1.9e+02;
Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGTCAGGCATCTAG 23
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Db 13217 TATAGTGTTCAGGCATCCAG 13239

RESULT 14
US-07-644-372-1/C
; Sequence 1, Application US/07644372
; Patent No. 5416009
; GENERAL INFORMATION:
; APPLICANT: Lazzeri, Mario E.
; APPLICANT: Nutman, Thomas B.
; APPLICANT: Weiss, Niklaus
; TITLE OF INVENTION: A DNA SEGMENT ENCODING A SPECIFIC
; OPERATING SYSTEM: IMMUNODIAGNOSTIC ANTIGEN
```



NUMBER OF SEQUENCES: 2  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: CUSHMAN, DARBY & CUSHMAN  
STREET: 1615 L. Street, N.W.  
CITY: Washington  
STATE: D.C.  
COUNTRY: USA  
ZIP: 20036  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/644,372  
FILING DATE: 19910123  
CLASSIFICATION: 435  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (202)861-3000  
TELEFAX: (202)822-0944  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 822 base pairs  
TYPE: NUCLEIC ACID  
STRANDEDNESS: double  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 52...507  
US-07-644-372-1

Query Match 61.7%; Score 14.8; DB 1; Length 822;  
Best Local Similarity 88.9%; Pred. No. 1.6e+02;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
Qy 1 TCTAGTGTCTGTCGAGCA 18  
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Db 689 TCTAGTGTCTGTCGAGCA 672

RESULT 15  
US-08-296-014A-3  
Sequence 3, Application US/08296014A  
Patent No. 5716834  
GENERAL INFORMATION:  
APPLICANT: Ding, Jeak Ling  
APPLICANT: Ho, Bow  
TITLE OF INVENTION: The Cloned Factor C cDNA of the  
TITLE OF INVENTION: Singapore Horseshoe Crab, Carcinoscorpius  
TITLE OF INVENTION: rotundicauda and Purification of Factor C proenzyme  
NUMBER OF SEQUENCES: 4  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Birch, Stewart, Kolasch & Birch  
STREET: 8110 Gatehouse Road, Suite 500 East  
CITY: Falls Church  
STATE: Virginia  
COUNTRY: USA  
ZIP: 22042  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/296,014A  
FILING DATE:  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Murphy, Jr., Gerald M.  
REGISTRATION NUMBER: 28,977  
REFERENCE/DOCKET NUMBER: 1781-105P  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (703) 203-8000

TELEFAX: (703) 203-8050  
TELEX: 248345  
INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3448 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: both  
MOLECULE TYPE: cDNA  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Carcinoscorpius rotundicauda  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 18..3074  
US-08-296-014A-3

Query Match 61.7%; Score 14.8; DB 1; Length 3448;  
Best Local Similarity 88.9%; Pred. No. 2e+02;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
Qy 5 GTGTCGTGTCGAGGCATCTA 22  
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Db 1136 GTGTCGTGTCGAGGCATCCA 1153

Search completed: January 4, 2003, 00:11:20  
Job time : 81.1009 secs



GenCore version 5.1.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:41 ; Search time 8.10095 Seconds  
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Title: US-09-787-562-11

Perfect score: 24

Sequence: 1 tctagtgtgtgcaggcatctagt 24

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 381593 seqs, 216252194 residues

Total number of hits satisfying chosen parameters: 763186

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Published Applications\_NA.\*  
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14: /cgn2\_6/ptodata/1/pubpna/US60\_PUBCOMB.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
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C 2	16.6	69.2	2056	12	US-10-003-152-23
C 3	16.2	67.5	371	10	US-09-960-352-2882
C 4	16.2	67.5	557	10	US-09-974-300-4795
C 5	16	66.7	159	10	US-09-864-761-22977
C 6	16	66.7	387	10	US-09-878-574-2017
C 7	16	66.7	456	10	US-09-864-761-6257
C 8	16	66.7	1005	10	US-09-815-242-6973
C 9	15.8	65.8	88191	10	US-09-799-799-3
C 10	15.6	65.0	200	10	US-09-864-761-24121
C 11	15.6	65.0	551	10	US-09-864-761-7458
C 12	15.6	65.0	1845	9	US-09-738-626-3487
C 13	15.6	65.0	40392	10	US-09-954-456-44
C 14	15.6	65.0	40392	10	US-09-954-456-44
C 15	15.4	64.2	598	9	US-10-001-835-62
C 16	15.2	63.3	299	10	US-09-294-093B-3520
C 17	15.2	63.3	684973	10	US-09-263-959-1
C 18	15	62.5	419	9	US-09-796-692-7030
C 19	15	62.5	594	10	US-09-879-336-155

C 20	15	62.5	954	10	US-09-815-242-9540	Sequence 9540, Ap
C 21	15	62.5	1197	9	US-09-938-842A-793	Sequence 793, App
C 22	15	62.5	2141	10	US-09-917-800A-1511	Sequence 1511, Ap
C 23	15	62.5	2256	10	US-09-899-471-1	Sequence 1, Appli
C 24	15	62.5	2328	10	US-09-899-471-4	Sequence 4, Appli
C 25	15	62.5	2538	9	US-10-098-841-313	Sequence 313, App
C 26	15	62.5	17056	10	US-09-893-238-3	Sequence 3, Appli
C 27	15	62.5	167343	10	US-09-962-436-281	Sequence 281, App
C 28	15	62.5	167343	10	US-09-964-824A-273	Sequence 273, App
C 29	15	62.5	174424	10	US-09-967-768A-314	Sequence 314, App
C 30	15	62.5	196285	10	US-09-880-107-3814	Sequence 3814, Ap
C 31	14.8	61.7	217	10	US-09-867-701-4514	Sequence 4514, Ap
C 32	14.8	61.7	580	10	US-09-864-761-8242	Sequence 8242, Ap
C 33	14.8	61.7	4369	10	US-09-769-097-1	Sequence 1, Appli
C 34	14.8	61.7	4425	10	US-09-769-097-3	Sequence 3, Appli
C 35	14.8	61.7	15535	10	US-09-764-877-2855	Sequence 2855, Ap
C 36	14.8	61.7	203654	10	US-09-820-905-3	Sequence 3, Appli
C 37	14.6	60.8	127	10	US-09-867-701-10523	Sequence 10523, A
C 38	14.6	60.8	192	10	US-09-867-701-8124	Sequence 8124, Ap
C 39	14.6	60.8	196	10	US-09-864-761-30975	Sequence 30975, A
C 40	14.6	60.8	374	9	US-09-886-242A-3	Sequence 3, Appli
C 41	14.6	60.8	374	9	US-10-027-603-3	Sequence 3, Appli
C 42	14.6	60.8	387	10	US-09-983-965-3685	Sequence 3685, Ap
C 43	14.6	60.8	429	9	US-09-796-692-8460	Sequence 8460, Ap
C 44	14.6	60.8	457	10	US-09-864-761-14419	Sequence 14419, A
C 45	14.6	60.8	461	10	US-09-864-761-15009	Sequence 15009, A

## ALIGNMENTS

RESULT 1  
US-10-003-152-15/c  
; Sequence 15, Application US/10003152  
; Patent No. US20020151494A1  
; GENERAL INFORMATION:  
; APPLICANT: Shimkets, Richard  
; APPLICANT: Fernandes, Elma  
; APPLICANT: Vernet, Corine  
; APPLICANT: Yang, Meljia  
; APPLICANT: Boldog, Ferenc  
; APPLICANT: Herimann, John  
; TITLE OF INVENTION: No. US20020151494A1el Amino Acid Sequences for Human Semaphori  
; FILE REFERENCE: 15966-554 Cura-54 CON-S12  
; CURRENT APPLICATION NUMBER: US/10/003.152  
; CURRENT FILING DATE: 2001-11-02  
; PRIOR APPLICATION NUMBER: 09/604,286  
; PRIOR FILING DATE: 2000-06-22  
; PRIOR APPLICATION NUMBER: 60/140,584  
; PRIOR FILING DATE: 1999-06-23  
; NUMBER OF SEQ ID NOS: 49  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 15  
; LENGTH: 1930  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: (148)..(1755)  
; NAME/KEY: variation  
; LOCATION: (1)..(1930)  
; OTHER INFORMATION: N may be any nucleotide  
US-10-003-152-15

Query Match 69.2%; Score 16.6; DB 12; Length 1930;  
Best Local Similarity 82.6%; Pred. No. 36;  
Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 2 CTAGTGTCTGTCAGGCATCTAGT 24  
Db 284 CTAGTCTCTGTCAGGCATCTCGT 262

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RESULT 2
US-10-003-152-23/C
; Sequence 23, Application US/10003152
; Patent No. US20020151494A1
; GENERAL INFORMATION:
; APPLICANT: Shimkets, Richard
; APPLICANT: Fernandes, Elma
; APPLICANT: Vernet, Corine
; APPLICANT: Yang, Meijia
; APPLICANT: Boldog, Ferenc
; APPLICANT: Herrmann, John
; TITLE OF INVENTION: NO. US20020151494A1el Amino Acid Sequences for Human Semaphorin-1
; FILE REFERENCE: 15966-554 Cura-54 CON-S12
; CURRENT APPLICATION NUMBER: US/10/003,152
; CURRENT FILING DATE: 2001-11-02
; PRIOR APPLICATION NUMBER: 09/604,286
; PRIOR FILING DATE: 2000-06-22
; PRIOR APPLICATION NUMBER: 60/140,584
; PRIOR FILING DATE: 1999-06-23
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 23
; LENGTH: 2056
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (7)..(1608)
; NAME/KEY: variation
; LOCATION: (1)..(2056)
; OTHER INFORMATION: N may be any nucleotide
; NAME/KEY: variation
; LOCATION: (1)..(2056)
; OTHER INFORMATION: N may be any nucleotide
US-10-003-152-23

Query Match 69.2%; Score 16.6; DB 12; Length 2056;
Best Local Similarity 82.6%; Pred. No. 36;
Matches 19; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 CTAGTGTGTCGAGGCATCTAGT 24
||||| ||||||| |||||||
Db 137 CTAGTCTCTGTCGAGGCATCTCGT 115

RESULT 3
US-09-960-352-2882
; Sequence 2882, Application US/09960352
; Patent No. US20020137139A1
; GENERAL INFORMATION:
; APPLICANT: Warren, Wesley C.
; APPLICANT: Tao, Nengbing
; APPLICANT: Byatt, John C.
; APPLICANT: Mathialagan, Nagappan
; TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH LACTATION AND
; FILE REFERENCE: 16511.006/37-21(10298)C
; CURRENT APPLICATION NUMBER: US/09/960,352
; CURRENT FILING DATE: 2001-09-24
; NUMBER OF SEQ ID NOS: 15112
; SEQ ID NO 2882
; LENGTH: 371
; TYPE: DNA
; ORGANISM: Bos taurus
; OTHER INFORMATION: Clone ID: 13-LIB3057-007-01-K1-D1
US-09-960-352-2882

Query Match 67.5%; Score 16.2; DB 10; Length 371;
Best Local Similarity 85.7%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TAGTGTGTCGAGGCATCTAGT 23
| ||||| ||||||| |||||||
```

```
Db 224 TTGTGTCATCGAGGCATCTAG 244

RESULT 4
US-09-974-300-4795
; Sequence 4795, Application US/09974300
; Patent No. US20020146721A1
; GENERAL INFORMATION:
; APPLICANT: Berka, Randy M.
; APPLICANT: Clausen, Ib Groth
; TITLE OF INVENTION: Methods for Monitoring Multiple Gene
; FILE REFERENCE: 10085.500-US
; CURRENT APPLICATION NUMBER: US/09/974,300
; CURRENT FILING DATE: 2001-10-05
; PRIOR APPLICATION NUMBER: 09/680,598
; PRIOR FILING DATE: 2000-10-06
; PRIOR APPLICATION NUMBER: 60/279,526
; PRIOR FILING DATE: 2001-03-27
; NUMBER OF SEQ ID NOS: 8481
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 4795
; LENGTH: 557
; TYPE: DNA
; ORGANISM: Bacillus clausii
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)..(557)
; OTHER INFORMATION: n = A,T,C or G
US-09-974-300-4795

Query Match 67.5%; Score 16.2; DB 10; Length 557;
Best Local Similarity 85.7%; Pred. No. 52;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CTAGTGTGTCGAGGCATCTA 22
|| ||||| |||| |||||||
Db 292 CTCGTGTCCTGCAAGCATCTA 312

RESULT 5
US-09-864-761-22977/c
; Sequence 22977, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharron G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO
; FILE REFERENCE: Aeomica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263.6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
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;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00668  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00663  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00662  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00661  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00670  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: US 60/234,687  
;; PRIOR FILING DATE: 2000-09-21  
;; PRIOR APPLICATION NUMBER: US 09/608,408  
;; PRIOR FILING DATE: 2000-06-30  
;; PRIOR APPLICATION NUMBER: US 09/774,203  
;; PRIOR FILING DATE: 2001-01-29  
;; NUMBER OF SEQ ID NOS: 49117  
;; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
;; SEQ ID NO 22977  
;; LENGTH: 159  
;; TYPE: DNA  
;; ORGANISM: Homo sapiens  
;; FEATURE:  
;; OTHER INFORMATION: MAP TO AC004549.1  
;; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 15  
;; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 13  
;; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 15  
;; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 14  
;; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 15  
;; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 17  
;; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 14  
;; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 15  
;; OTHER INFORMATION: NT HIT: AFL35399.1, EVALUATE 3.60e-01  
;; OTHER INFORMATION: EST\_HUMAN HIT: AI833003.1, EVALUATE 2.00e-77  
US-09-864-761-22977

Query Match 66.7%; Score 16; DB 10; Length 159;  
Best Local Similarity 79.2%; Pred. No. 60;  
Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 TCTAGTGTGTCGACGCGATCTAGT 24  
||| |||| |||| |||| ||||  
DB 158 TCTCTGTGTCGACGCGCGAGT 135

RESULT 6  
US-09-878-574-2017  
;; Sequence 2017, Application US/09878574  
;; Patent No. US20020110548A1  
;; GENERAL INFORMATION:  
;; APPLICANT: Byrum, Joseph R.  
;; APPLICANT: La Rosa, Thomas J.  
;; APPLICANT: Thompson, Michael D.  
;; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated with  
;; TITLE OF INVENTION: Plants  
;; FILE REFERENCE: 38-21(15401)B  
;; CURRENT APPLICATION NUMBER: US/09/878,574  
;; CURRENT FILING DATE: 2001-12-21  
;; PRIOR APPLICATION NUMBER: 09/333,535  
;; PRIOR FILING DATE: 1999-06-14  
;; NUMBER OF SEQ ID NOS: 15775  
;; SEQ ID NO 2017  
;; LENGTH: 387  
;; TYPE: DNA  
;; ORGANISM: Glycine max  
;; OTHER INFORMATION: Clone ID: LIB3028-030-Q1-B1-C4  
US-09-878-574-2017

Query Match 66.7%; Score 16; DB 10; Length 387;  
Best Local Similarity 79.2%; Pred. No. 63;  
Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 TCTAGTGTGTCGACGCGATCTAGT 24  
||| |||| |||| |||| ||||  
DB 258 TCTACACTCTTGCAGCCATCTAGT 281

RESULT 7  
US-09-864-761-6257/c  
;; Sequence 6257, Application US/09864761  
;; Patent No. US20020048763A1  
;; GENERAL INFORMATION:  
;; APPLICANT: Penn, Sharron G.  
;; APPLICANT: Rank, David R.  
;; APPLICANT: Hanzel, David K.  
;; APPLICANT: Chen, Wensheng  
;; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
;; TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY  
;; FILE REFERENCE: Acomica-X-1  
;; CURRENT APPLICATION NUMBER: US/09/864,761  
;; CURRENT FILING DATE: 2001-05-23  
;; PRIOR APPLICATION NUMBER: US 60/180,312  
;; PRIOR FILING DATE: 2000-02-04  
;; PRIOR APPLICATION NUMBER: US 60/207,456  
;; PRIOR FILING DATE: 2000-05-26  
;; PRIOR APPLICATION NUMBER: US 09/632,366  
;; PRIOR FILING DATE: 2000-08-03  
;; PRIOR APPLICATION NUMBER: GB 24263.6  
;; PRIOR FILING DATE: 2000-10-04  
;; PRIOR APPLICATION NUMBER: US 60/236,359  
;; PRIOR FILING DATE: 2000-09-27  
;; PRIOR APPLICATION NUMBER: PCT/US01/00666  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00667  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00664  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00669  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00665  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00668  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00663  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00662  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00661  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: PCT/US01/00670  
;; PRIOR FILING DATE: 2001-01-30  
;; PRIOR APPLICATION NUMBER: US 60/234,687  
;; PRIOR FILING DATE: 2000-09-21  
;; PRIOR APPLICATION NUMBER: US 09/608,408  
;; PRIOR FILING DATE: 2000-06-30  
;; PRIOR APPLICATION NUMBER: US 09/774,203  
;; PRIOR FILING DATE: 2001-01-29  
;; NUMBER OF SEQ ID NOS: 49117  
;; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
;; SEQ ID NO 6257  
;; LENGTH: 456  
;; TYPE: DNA  
;; ORGANISM: Homo sapiens  
;; FEATURE:  
;; OTHER INFORMATION: MAP TO AC004549.1  
;; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 15  
;; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 13  
;; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 15  
;; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 14  
;; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 15  
;; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 17  
;; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 14  
;; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 15  
US-09-864-761-6257

Query Match 66.7%; Score 16; DB 10; Length 456;  
Best Local Similarity 79.2%; Pred. No. 64;  
Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGCAGGCATCTAGT 24  
||| |||| |||| |||| |||| |||| |||| ||||  
DB 435 TCTCTTGTCTGCAGGCAGCGAGT 412

## RESULT 8

US-09-815-242-6973  
; Sequence 6973, Application US/09815242  
; Patent No. US20020061569A1  
; GENERAL INFORMATION:  
; APPLICANT: Haselbeck, Robert  
; APPLICANT: Ohlsen, Karl L.  
; APPLICANT: Zyskind, Judith W.  
; APPLICANT: Wall, Daniel  
; APPLICANT: Trawick, John D.  
; APPLICANT: Carr, Grant J.  
; APPLICANT: Yamamoto, Robert T.  
; APPLICANT: Xu, H. Howard

; TITLE OF INVENTION: Identification of Essential Genes in

; FILE REFERENCE: ELITRA.011A

; CURRENT APPLICATION NUMBER: US/09/815,242

; CURRENT FILING DATE: 2001-03-21

; PRIOR APPLICATION NUMBER: 60/191,078

; PRIOR FILING DATE: 2000-03-21

; PRIOR APPLICATION NUMBER: 60/206,848

; PRIOR FILING DATE: 2000-05-23

; PRIOR APPLICATION NUMBER: 60/207,727

; PRIOR FILING DATE: 2000-05-26

; PRIOR APPLICATION NUMBER: 60/242,578

; PRIOR FILING DATE: 2000-10-23

; PRIOR APPLICATION NUMBER: 60/253,625

; PRIOR FILING DATE: 2000-11-27

; PRIOR APPLICATION NUMBER: 60/257,931

; PRIOR FILING DATE: 2000-12-22

; PRIOR APPLICATION NUMBER: 60/269,308

; PRIOR FILING DATE: 2001-02-16

; NUMBER OF SEQ ID NOS: 14110

; SOFTWARE: FastSeq for Windows Version 4.0

; SEQ ID NO 6973

; LENGTH: 1005

; TYPE: DNA

; ORGANISM: Haemophilus influenzae

; FEATURE:

; NAME/KEY: CDS

; LOCATION: (1)...(1005)

US-09-815-242-6973

Query Match 66.7%; Score 16; DB 10; Length 1005;  
Best Local Similarity 79.2%; Pred. No. 67;  
Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 TCTAGTGTCTGCAGGCATCTAGT 24  
||| |||| |||| |||| |||| |||| |||| ||||  
DB 19 TTAGTGGCGTGCAGCCCTCTGGT 42

## RESULT 9

US-09-799-799-3  
; Sequence 3, Application US/09799799  
; Patent No. US20020132291A1  
; GENERAL INFORMATION:

; APPLICANT: YE, Jane et al.

; TITLE OF INVENTION: ISOLATED HUMAN RAS-LIKE PROTEINS,

; TITLE OF INVENTION: NUCLEIC ACID MOLECULES ENCODING THESE HUMAN RAS-LIKE

; TITLE OF INVENTION: PROTEINS, AND USES THEREOF

; FILE REFERENCE: C1001157

; CURRENT APPLICATION NUMBER: US/09/799,799

; CURRENT FILING DATE: 2001-03-07

; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: FastSeq for Windows Version 4.0

; SEQ ID NO 3

; LENGTH: 88191

; TYPE: DNA

; ORGANISM: Human

; FEATURE:

; NAME/KEY: misc\_feature

; LOCATION: (1)...(88191)

; OTHER INFORMATION: n = A,T,C or G

US-09-799-799-3

Query Match 65.8%; Score 15.8; DB 10; Length 88191;  
Best Local Similarity 89.5%; Pred. No. 1.1e+02;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 6 TGTCTGTCAGGCATCTAGT 24  
|||| |||| |||| |||| |||| |||| |||| ||||  
DB 72640 TGTCTGTCAGGCATCTAGT 72658

## RESULT 10

US-09-864-761-24121

; Sequence 24121, Application US/09864761

; Patent No. US20020048763A1

; GENERAL INFORMATION:

; APPLICANT: Penn, Sharron G.

; APPLICANT: Rank, David R.

; APPLICANT: Hanzel, David K.

; APPLICANT: Chen, Wensheng

; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO

; TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY

; FILE REFERENCE: Aecmca-X-1

; CURRENT APPLICATION NUMBER: US/09/864,761

; CURRENT FILING DATE: 2001-05-23

; PRIOR APPLICATION NUMBER: US 60/180,312

; PRIOR FILING DATE: 2000-02-04

; PRIOR APPLICATION NUMBER: US 60/207,456

; PRIOR FILING DATE: 2000-05-26

; PRIOR APPLICATION NUMBER: US 09/632,366

; PRIOR FILING DATE: 2000-08-03

; PRIOR APPLICATION NUMBER: GB 24263.6

; PRIOR FILING DATE: 2000-10-04

; PRIOR APPLICATION NUMBER: US 60/236,359

; PRIOR FILING DATE: 2000-09-27

; PRIOR APPLICATION NUMBER: PCT/US01/00666

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00667

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00664

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00669

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00665

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00668

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00663

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00662

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00661

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00670

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: US 60/234,687

; PRIOR FILING DATE: 2000-09-21

; PRIOR APPLICATION NUMBER: US 09/608,408

; PRIOR FILING DATE: 2000-06-30

; PRIOR APPLICATION NUMBER: US 09/774,203

; PRIOR FILING DATE: 2001-01-29

; NUMBER OF SEQ ID NOS: 49117

; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1

; SEQ ID NO 24121  
; LENGTH: 200  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: MAP TO AL139347.2  
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.1  
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3  
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2  
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.88  
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1  
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1  
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.3  
; OTHER INFORMATION: EST\_HUMAN HIT: BF690757.1, EVALUE 3.00e-49  
; OTHER INFORMATION: NT HIT: g14758789, EVALUE 2.00e-52  
; OTHER INFORMATION: SWISSPROT HIT: O43920, EVALUE 3.00e-18  
US-09-864-761-24121

Query Match 65.0%; Score 15.6; DB 10; Length 200;  
Best Local Similarity 81.8%; Pred. No. 95;  
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 3 TAGTGTCTGCGAGGCATCTAGT 24  
| ||||| ||||| ||||| ||  
Db 45 TGGTGTGCGAGGAACTCTTGT 66

RESULT 11  
US-09-864-761-7458  
; Sequence 7458, Application US/09864761  
; Patent No. US20020048763A1  
; GENERAL INFORMATION:  
; APPLICANT: Penn, Sharron G.  
; APPLICANT: Rank, David R.  
; APPLICANT: Hanzel, David K.  
; APPLICANT: Chen, Wensheng  
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
; FILE REFERENCE: Aomica-X-1  
; CURRENT APPLICATION NUMBER: US/09/864,761  
; PRIOR FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/180,312  
; PRIOR FILING DATE: 2000-02-04  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 09/632,366  
; PRIOR FILING DATE: 2000-08-03  
; PRIOR APPLICATION NUMBER: GB 24263.6  
; PRIOR FILING DATE: 2000-10-04  
; PRIOR APPLICATION NUMBER: US 60/236,359  
; PRIOR FILING DATE: 2000-09-27  
; PRIOR APPLICATION NUMBER: PCT/US01/00666  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00662  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00661  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00670  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 60/234,687  
; PRIOR FILING DATE: 2000-09-21

; PRIOR APPLICATION NUMBER: US 09/608,408  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: US 09/774,203  
; PRIOR FILING DATE: 2001-01-29  
; NUMBER OF SEQ ID NOS: 49117  
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
; SEQ ID NO 7458  
; LENGTH: 551  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: MAP TO AL139347.2  
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.1  
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3  
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2  
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.88  
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1  
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1  
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.3  
US-09-864-761-7458

Query Match 65.0%; Score 15.6; DB 10; Length 551;  
Best Local Similarity 81.8%; Pred. No. 1e+02;  
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 3 TAGTGTCTGCGAGGCATCTAGT 24  
| ||||| ||||| ||||| ||  
Db 152 TGGTGTGCGAGGAACTCTTGT 173

RESULT 12  
US-09-738-626-3487  
; Sequence 3487, Application US/09738626  
; Publication No. US20020197605A1  
; GENERAL INFORMATION:  
; APPLICANT: NAKAGAWA, SATOSHI  
; APPLICANT: MIZOGUCHI, HIROSHI  
; APPLICANT: ANDO, SEIKO  
; APPLICANT: HAYASHI, MIKIRO  
; APPLICANT: OCHIALI, KEIKO  
; APPLICANT: YOKOI, HARUHIKO  
; APPLICANT: TATEISHI, NAKO  
; APPLICANT: SENOH, AKIHIRO  
; APPLICANT: IKEDA, MASATO  
; APPLICANT: OZAKI, AKIO  
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES  
; FILE REFERENCE: 249-125  
; CURRENT APPLICATION NUMBER: US/09/738,626  
; CURRENT FILING DATE: 2000-12-18  
; PRIOR APPLICATION NUMBER: JP 99/377484  
; PRIOR FILING DATE: 1999-12-16  
; PRIOR APPLICATION NUMBER: JP 00/159162  
; PRIOR FILING DATE: 2000-04-07  
; PRIOR APPLICATION NUMBER: JP 00/280988  
; PRIOR FILING DATE: 2000-08-03  
; NUMBER OF SEQ ID NOS: 7059  
; SOFTWARE: PatentIn ver. 3.0  
; SEQ ID NO 3487  
; LENGTH: 1845  
; TYPE: DNA  
; ORGANISM: Corynebacterium glutamicum  
US-09-738-626-3487

Query Match 65.0%; Score 15.6; DB 9; Length 1845;  
Best Local Similarity 81.8%; Pred. No. 1.1e+02;  
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 TCTAGTGTCTGCGAGGCATCTA 22  
| | | | | | | | | | | | | | | | | |  
Db 1133 TCTTTGGTCTGCGAGGCATCGA 1154

RESULT 13

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US-09-954-456-44
; Sequence 44, Application US/09954456
; Patent No. US20020115057A1
; GENERAL INFORMATION:
; APPLICANT: Young, Paul
; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using Cand
; FILE REFERENCE: 689290-76
; CURRENT APPLICATION NUMBER: US/09/954,456
; PRIOR FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/60/233,617
; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/234,923
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,840
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,863
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 44
; LENGTH: 40392
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-44

Query Match 65.0%; Score 15.6; DB 10; Length 40392;
Best Local Similarity 81.8%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 3 TAGTGTCTGTCAGGCATCTAGT 24
|| |||| |||| |||| |||| ||
Db 27507 TATTCTCTGCAAGCATCTGGT 27528

RESULT 14
US-09-954-456-687
; Sequence 687, Application US/09954456
; Patent No. US20020115057A1
; GENERAL INFORMATION:
; APPLICANT: Young, Paul
; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using Cand
; FILE REFERENCE: 689290-76
; CURRENT APPLICATION NUMBER: US/09/954,456
; PRIOR FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/60/233,617
; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 44
; LENGTH: 40392
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-44

Query Match 65.0%; Score 15.6; DB 10; Length 40392;
Best Local Similarity 81.8%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 3 TAGTGTCTGTCAGGCATCTAGT 24
|| |||| |||| |||| |||| ||
Db 27507 TATTCTCTGCAAGCATCTGGT 27528

RESULT 14
US-09-954-456-687
; Sequence 687, Application US/09954456
; Patent No. US20020115057A1
; GENERAL INFORMATION:
; APPLICANT: Young, Paul
; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using Cand
; FILE REFERENCE: 689290-76
; CURRENT APPLICATION NUMBER: US/09/954,456
; PRIOR FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/60/233,617
; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
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US-09-954-456-687
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,840
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,863
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 687
; LENGTH: 40392
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-687

Query Match 65.0%; Score 15.6; DB 10; Length 40392;
Best Local Similarity 81.8%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 3 TAGTGTCTGTCAGGCATCTAGT 24
|| |||| |||| |||| |||| ||
Db 27507 TATTGTCTGCAAGCATCTGGT 27528

RESULT 15
US-10-001-835-62/c
; Sequence 62, Application US/10001835
; Patent No. US20020160387A1
; GENERAL INFORMATION:
; APPLICANT: Salceda, Susana
; APPLICANT: Macina, Roberto
; APPLICANT: Recipon, Herve
; APPLICANT: Cafferkey, Robert
; APPLICANT: Sun, Yongming
; APPLICANT: Liu, Chenghua
; TITLE OF INVENTION: Compositions and Methods Relating to Ovary Specific Genes and
; FILE REFERENCE: DEX-0277
; CURRENT APPLICATION NUMBER: US/10/001,835
; CURRENT FILING DATE: 2001-11-20
; PRIOR APPLICATION NUMBER: 60/249,997
; PRIOR FILING DATE: 2000-11-20
; NUMBER OF SEQ ID NOS: 228
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 62
; LENGTH: 598
; TYPE: DNA
; ORGANISM: Homo sapien
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (270)..(270)
; OTHER INFORMATION: a, c, g or t
; NAME/KEY: misc_feature
; LOCATION: (330)..(330)
; OTHER INFORMATION: a, c, g or t
; NAME/KEY: misc_feature
; LOCATION: (332)..(332)
; OTHER INFORMATION: a, c, g or t
; NAME/KEY: misc_feature
; LOCATION: (334)..(334)
; OTHER INFORMATION: a, c, g or t
; NAME/KEY: misc_feature
; LOCATION: (533)..(533)
; OTHER INFORMATION: a, c, g or t
US-10-001-835-62

Query Match 64.2%; Score 15.4; DB 9; Length 598;
Best Local Similarity 94.1%; Pred. No. 1.3e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 8 TCGTGCAGGCATCTAGT 24
|| || |||| |||| |||| ||
Db 59 TCGAGCAGGCATCTAGT 43

Search completed: January 4, 2003, 01:06:35
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GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:52:46 ; Search time 2134.5 Seconds  
(without alignments) 3231.380 Million cell updates/sec

Title: US-09-787-562-9  
Perfect score: 237  
Sequence: 1 gctagagtcgtgcaggacgt.....cgaqccaccttcagcctctg 237

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues  
Total number of hits satisfying chosen parameters: 4109280

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000
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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : GenEmbl : \*

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2: gb\_ba:\*  
3: gb\_htg:\*  
4: gb\_in:\*  
5: gb\_om:\*  
6: gb\_ov:\*  
7: gb\_pat:\*  
8: gb\_ph:\*  
9: gb\_pl:\*  
10: gb\_pr:\*  
11: gb\_ro:\*  
12: gb\_sy:\*  
13: gb\_un:\*  
14: gb\_vl:\*  
15: em\_ba:\*  
16: em\_fun:\*  
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24: em\_ph:\*  
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26: em\_ro:\*  
27: em\_sts:\*  
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29: em\_vl:\*  
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33: em\_htg\_mus:\*  
34: em\_htg\_pln:\*  
35: em\_htg\_rod:\*  
36: em\_htg\_man:\*  
37: em\_htg\_vrt:\*  
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39: em\_htg\_hum:\*  
40: em\_htg\_mus:\*  
41: em\_htg\_inv:\*  
42: em\_htg\_inv:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Query			DB	ID	Description
	Score	Match	Length			
1	237	100.0	237	6	AX023667	Sequence
2	221	93.2	243	6	AX018563	Sequence
3	221	93.2	243	6	AX018639	Sequence
4	221	93.2	243	6	AX023661	Sequence
5	175	73.8	242	6	AX023682	Sequence
6	171	72.2	4219	12	RVU57027	Sequence
7	171	72.2	4465	12	RVU57025	Reporter vec
8	171	72.2	4869	12	CVU89940	Reporter vec
9	171	72.2	5010	6	AX339210	Cloning vec
10	171	72.2	5010	12	CVU47298	Cloning vec
11	171	72.2	5115	12	CVU89938	Sequence
12	171	72.2	5256	6	AX339208	Sequence
13	171	72.2	5256	12	CVU47296	Cloning vec
14	160	67.5	6320	12	AB038600	Sequence
15	157.2	66.3	5917	12	AB037684	Cloning vec
16	152.2	64.2	267	6	AX023685	Sequence
17	148.4	62.6	204	6	AX023687	Sequence
18	148	62.4	223	6	AX023683	Sequence
19	147.4	62.2	356	6	A58090	Sequence 1
20	147.4	62.2	356	6	AR095774	Sequence
21	146.8	61.9	5789	12	CVPLG2PRO	Cloning vec
22	146.8	61.9	5991	12	CVU09663	Cloning vec
23	146.8	61.9	6046	12	CVPLG2CON	Cloning vec
24	146.8	61.9	6248	12	CVU09661	Cloning vec
25	146.8	61.9	7674	12	CVU13187	Cloning vec
26	146.8	61.9	7931	12	CVU13186	Cloning vec
27	146.4	61.8	4506	12	CVPCATPRO	Cloning vec
28	146.4	61.8	4750	12	CVPCATCON	Cloning vec
29	145.2	61.3	7731	6	A94048	Sequence 29
30	145.2	61.3	7731	6	A94061	Sequence 42
31	145.2	61.3	7731	6	AX011133	Sequence
32	145.2	61.3	7731	6	AX011146	Sequence
33	144.8	61.1	5894	12	AF334827	Sequence
34	144.4	60.9	8772	12	AF136442	Cloning v
35	143.6	60.6	110000	2	LMPLCHR34_08	Continuation (9 of
36	142.8	60.3	350	6	E00567	E00567 DNA sequenc
37	142.8	60.3	455	6	E01399	E01399 DNA sequenc
C 38	142.8	60.3	5003	12	SYNPSV2CAT	M77788 pSV2-Cat.c1
C 39	142.8	60.3	5840	12	NE1EXPVECB	L07041 pMNeo euka
C 40	142.8	60.3	5846	12	NE1EXPVECA	L07040 pMNeo euka
41	142.8	60.3	8068	6	A94046	Sequence 27
42	142.8	60.3	8068	6	A94054	Sequence 35
43	142.8	60.3	8068	6	AX011131	Sequence
44	142.8	60.3	8068	6	AX011139	Sequence
45	142.8	60.3	13254	6	AR038307	Sequence

## ALIGNMENTS

[illegible]

DNA linear PAT 15-SEP-2000

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source		1. .237	
		/organism="synthetic construct"	
		/db_xref="taxon:32630"	
		/note="OBhrell"	56 t
BASE COUNT		43 a	82 c 56 g 56 t
ORIGIN			
Query Match		100.0%;	Score 237; DB 6; Length 237;
Best Local Similarity		100.0%;	Pred. No. 1.6e-62;
Matches 237;		Conservative 0;	Mismatches 0; Indels 0; Gaps 0;
QY	1	GCTAGAGTCGTCAGGACGTCGACATCTAGTCGTGCGAGGACATCTAGTCGTGCGCAGGAC	60
Db	1	GCTAGAGTCGTCAGGACGTCGACATCTAGTCGTGCGAGGACATCTAGTCGTGCGCAGGAC	60
QY	61	GTGACAGCTAGCCGGCTCGAGATCTCGATCTCGATCTCAATTAGTCAGCAACCATAG	120
Db	61	GTGACAGCTAGCCGGCTCGAGATCTCGATCTCGATCTCAATTAGTCAGCAACCATAG	120
QY	121	TCCCGCCCTAACTCCGCCCATCCGCCCTAACTCCGCCCATGTCGCCCATCTCCGC	180
Db	121	TCCCGCCCTAACTCCGCCCATCCGCCCTAACTCCGCCCATGTCGCCCATCTCCGC	180
QY	181	CCATAGTCGCTGACTAATTTTTTTTATATGACAGGCGCGCCCTCGCCTCTG	237
Db	181	CCATAGTCGCTGACTAATTTTTTTTATATGACAGGCGCGCCCTCGCCTCTG	237
RESULT 2			
AX018563			
LOCUS		AX018563	243 bp DNA linear PAT 07-SEP-2000
DEFINITION		Sequence 57 from Patent WO9945127.	
ACCESSION		AX018563	
VERSION		AX018563.1	GI:10042701
KEYWORDS		synthetic construct.	
SOURCE		synthetic construct.	
ORGANISM		artificial sequences.	
REFERENCE		1 (bases 1 to 243)	
AUTHORS		Kingsman,S.M., Mitrophanous,K., Patterson,A.V., Stratford,I.J., Griffiths,L. and Kan,O.	
TITLE		Enhanced prodrug activation	
JOURNAL		Patent: WO 9945127-A 57 10-SEP-1999;	
		KINGSMAN SUSAN MARY (GB); MITROPHANOUS KYRIACOS (GB); PATTERSON ADAM VORN (GB); STRATFORD IAN JAMES (GB); GRIFFITHS LEIGH (GB); KAN ON (GB); OXFORD BIOMEDICA LTD (GB)	
FEATURES		Location/Qualifiers	
source		1. .243	
		/organism="synthetic construct"	
		/db_xref="taxon:32630"	
		/note="Promoter"	
BASE COUNT		45 a	83 c 58 g 57 t
ORIGIN			
Query Match		93.2%;	Score 221; DB 6; Length 243;
Best Local Similarity		97.5%;	Pred. No. 1.4e-57;
Matches 237;		Conservative 0;	Mismatches 0; Indels 6; Gaps 1;
QY	1	GCTAGAGTCGTCGAGGACGTCGACATCTAGTCGTGTCGTCAGG-----CATCTAGTCGTCG	54
Db	1	GCTAGAGTCGTCGAGGACGTCGACATCTAGTCGTGTCGTCGAGGACGTCGATCTAGTCGTCG	60
QY	55	CAGGACGTGACGCTAGCCGGCTCGAGATCTGCGATCTGCGATCTCAATTAGTCAGCAA	114
Db	61	CAGGACGTGACGCTAGCCGGCTCGAGATCTGCGATCTGCGATCTCAATTAGTCAGCAA	120
QY	115	CCATAGTCGCGCCCTAACTCCGCCCATCCGCCCTAACTCCGCCCATGTCGCCCATTT	174
Db	121	CCATAGTCGCGCCCTAACTCCGCCCATCCGCCCTAACTCCGCCCATGTCGCCCATTT	180
QY	175	CTCGGCCCATTCGCTGACTAATTTTTTTTATATGACAGGCGCGCCCTCGGCCT	234
Db	181	CTCGGCCCATTCGCTGACTAATTTTTTTTATATGACAGGCGCGCCCTCGGCCT	240
QY	235	CTG 237	
Db	241	CTG 243	
RESULT 4			
AX023661			
LOCUS		AX023661	243 bp DNA linear PAT 15-SEP-2000
DEFINITION		Sequence 3 from Patent WO0017371.	
ACCESSION		AX023661	
VERSION		AX023661.1	GI:10184022
KEYWORDS		synthetic construct.	
SOURCE		synthetic construct.	
ORGANISM		artificial sequences.	
REFERENCE		1 (bases 1 to 243)	
AUTHORS		Binley,K.M. and Naylor,S.	
TITLE		Polynucleotide constructs and uses thereof	
JOURNAL		Patent: WO 0017371-A 3 30-MAR-2000;	

BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD (GB)

## FEATURES

Source Location/Qualifiers

1..243  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="OBHrel"

BASE COUNT 45 a 83 c 58 g 57 t  
ORIGIN

Query Match 93.2%; Score 221; DB 6; Length 243;  
Best Local Similarity 97.5%; Pred. No. 1.4e-57;

Matches 237; Conservative 0; Mismatches 0; Indels 5; Gaps 1;

QY 1 GCTAGAGTCGTCAGAGAGTGACATCTAGTGTGCGTGCAGG-----CATCTAGTGTGCTG 54

Db 1 GCTAGAGTCGTCAGAGAGTGACATCTAGTGTGCGTGCAGAGCGTGACATCTAGTGTGCTG 60

QY 55 CAGGACGTGACAGCTAGAGCCGGGTCGAGATCTGCGATCTGCATCTCAATTAGTCAGCAA 114

Db 61 CAGGACGTGACAGCTAGAGCCGGGTCGAGATCTGCGATCTGCATCTCAATTAGTCAGCAA 120

QY 115 CCATAGTCCCGCCCTAAGTCCGCGCCATCCCGCCCTAACTCCGCCAGTTCCCGCCCAT 174

Db 121 CCATAGTCCCGCCCTAAGTCCGCGCCATCCCGCCCTAACTCCGCCAGTTCCCGCCCAT 180

QY 175 CTCGCCCGCCCTGCTGACTAATTTTTTATTATGACAGAGCGCGCGCTCGGCCT 234

Db 181 CTCGCCCGCCCTGCTGACTAATTTTTTATTATGACAGAGCGCGCGCTCGGCCT 240

QY 235 CTG 237  
Db 241 CTG 243

## RESULT 5

AX023682

LOCUS

DEFINITION

AX023682

VERSION

AX023682.1

KEYWORDS

SYNTHETIC CONSTRUCT.

ARTIFICIAL CONSTRUCT

ARTIFICIAL SEQUENCES.

REFERENCE

1 (bases 1 to 242)

AUTHORS

Binley,K.M. and Naylor,S.

TITLE

Polynucleotide constructs and uses thereof

JOURNAL

Patent: WO 0017371-A 24 30-MAR-2000;

BINLEY KATIE MARY (GB) ; NAYLOR STUART (GB) ; OXFORD BIOMEDICA LTD (GB)

FEATURES

Source Location/Qualifiers

1..242

/organism="synthetic construct"

/db\_xref="taxon:32630"

/note="Synthetic construct"

BASE COUNT 45 a 93 c 47 g 57 t

ORIGIN

Query Match 73.8%; Score 175; DB 6; Length 242;

Best Local Similarity 84.8%; Pred. No. 2.4e-43;

Matches 196; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

QY 7 GTCGTGACGACGTCGATCTAGTGTGCGTGCAGGATCTAGTGTGCGTGCAGGACGTGACA 66

Db 12 GTCGTGACGACGTCGATCTAGTGTGCGTGCAGGATCTAGTGTGCGTGCAGGACGTGACA 71

QY 67 GCTAGCCCGGGCTCGAGATCTGCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 126

Db 72 TCTAGCCCGGGCTCGAGATCTGCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 131

QY 127 CCTACTCGGCCCATCCCGCCCTAACTCCGCCAGTTCGCCGCCATCTCTCGGCCCATC 186

|||||

Db 132 CCCTAACTCCGCCCATCCCGCCCTAACTCCGCCAGTTCGCCCATCTCTCGGCCCATC 191

QY 187 GCTGACTAATTTTTTTTATTATGACAGAGCGCGAGCGCGCTCGGCTCTG 237

Db 192 GCTGACTAATTTTTTTTATTATGACAGAGCGCGAGCGCGCTCGGCTCTG 242

## RESULT 6

RVU57027

LOCUS

DEFINITION

RVU57027

ACCESSION

U57027.2

VERSION

U57027.2

KEYWORDS

ORGANISM

Reporter vector pCAT<R>3-Promoter vector.

ARTIFICIAL SEQUENCES; VECTORS.

REFERENCE

1 (bases 1 to 4219)

AUTHORS

Groskreutz,D.J., Vavra,S., Lesley,S. and Schenborn,E.

TITLE

CAT Reporter Systems: New pCAT(R)3 Reporter Vectors and Antibodies

JOURNAL

Promega Notes 55, 2-9 (1996)

REFERENCE

2 (bases 1 to 4219)

AUTHORS

Groskreutz,D.J. and Vavra,S.

TITLE

Direct Submission

JOURNAL

Submitted (30-APR-1996) Production, Promega Corporation, 5445 East

Cheryl Parkway, Madison, WI 53711, USA

REFERENCE

3 (bases 1 to 4219)

AUTHORS

Groskreutz,D.J. and Vavra,S.

TITLE

Direct Submission

JOURNAL

Submitted (24-APR-2001) Production, Promega Corporation, 5445 East

Cheryl Parkway, Madison, WI 53711, USA

REMARK

Sequence update by submitter

COMMENT

On Apr 24, 2001 this sequence version replaced gi:1399741.

FEATURES

Source Location/Qualifiers

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/db\_xref="taxon:47943"

misc\_feature

1..41

/note="multiple cloning site"

promoter

48..250

/note="SV40"

misc\_feature

185

/note="SV40 promoter directed transcriptional start site"

misc\_feature

191

/note="SV40 promoter directed transcriptional start site"

misc\_feature

196

/note="SV40 promoter directed transcriptional start site"

intron

295..427

/note="SV40 promoter directed transcriptional start site"

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/gene="CAT"

CDS

483..1142

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/codon\_start=1

/transl\_table=11

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/protein\_id="AAB39977.1"

/db\_xref="GI:1399742"

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ANMDFPAPVFTMGKYVTQGDVKVLMPLAIQVHVAVCDFHVRMLNELQQYCDWOGG

A"

1173..1394

/note="SV40 late polyA region"

complement(1462..1481)

/note="RV primer 4 sequencing primer binding site"

1719

/note="ColEI-derived plasmid replication origin"

complement(2481..3341)

/gene="Ampr"

CDS

complement(2481..3341)

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/ gene="AmpR"
/ function="ampicillin resistance"
/ codon_start=1
/ transl_table=11
/ product="beta-lactamase"
/ protein_id="AAB39978.1"
/ db_xref="GI:1399743"
/ translation="MSIQHFRVALIPFFAAFLPVPFAHPETLVKVKDAEDQLGARVGY
IELDLNSGKILSEFRPEERPMSTFKVLLCGAVLSRIDAGOEQLGRRIRHYQNDLVE
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DRWPELNEAIPNDERDTPMPVAMATTIRKLITGLTLLASRQQLIDNMEADKVGAPL
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EIGASLIKHW"
3474..3928
/ note="fl origin"
/ note="upstream"
/ note="RV primer 3 sequencing primer binding site"
BASE COUNT 1060 a 1046 c 992 g 1121 t
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Best Local Similarity 100.0%; Pred. No. 5.5e-42;
Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 67 GCTAGCCGGGCTCGAGATCTCGCATCTCAATTAGTCAGCAACCATAGTCCCGC 126
Db 21 GCTAGCCGGGCTCGAGATCTCGCATCTCAATTAGTCAGCAACCATAGTCCCGC 80
Qy 127 CCTAACTCCGCCCATCCGCCCTCACTCGCCAGTTCGCCCATCTCCGCCCATC 186
Db 81 CCTAACTCCGCCCATCCGCCCTCACTCGCCAGTTCGCCCATCTCCGCCCATC 140
Qy 187 GCTGACTAATTTTTTTTATTTATTCAGAGCGCGAGCGCGCTCGGCTCTG 237
Db 141 GCTGACTAATTTTTTTTATTTATTCAGAGCGCGAGCGCGCTCGGCTCTG 191

RESULT 7
RVU57025 4465 bp DNA linear SYN 24-APR-2001
LOCUS Reporter vector pCAT<R>3-Control vector, complete sequence.
DEFINITION U57025
ACCESSION U57025
VERSION U57025.2 GI:13775606
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS Groskreutz,D.J., Vavra,S., Lesley,S. and Schenborn,E.
TITLE CAT Reporter Systems: New pCAT(R)3 Reporter Vectors and Antibodies
Provide Increased Expression and Detection Capabilities
JOURNAL Promega Notes 55, 2-9 (1996)
REFERENCE 2 (bases 1 to 4465)
AUTHORS Groskreutz,D.J. and Vavra,S.
TITLE Direct Submission
JOURNAL Submitted (30-APR-1996) Production, Promega Corporation, 5445 East
Ceryl Parkway, Madison, WI 53711, USA
REFERENCE 3 (bases 1 to 4465)
AUTHORS Groskreutz,D.J. and Vavra,S.
TITLE Direct Submission
JOURNAL Submitted (24-APR-2001) Production, Promega Corporation, 5445 East
Ceryl Parkway, Madison, WI 53711, USA
REMARK Sequence update by submitter
COMMENT On Apr 24, 2001 this sequence version replaced gi:1399735.
FEATURES
source
1..4465
/organism="Reporter vector pCAT<R>3-Control vector"
/db_xref="taxon:47940"
misc_feature 1..41
/ note="multiple cloning site"
promoter 48..250

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/ note="SV40 promoter"
185
/ note="SV40 promoter directed transcriptional start site"
191
/ note="SV40 promoter directed transcriptional start site"
196
/ note="SV40 promoter directed transcriptional start site"
295..427
/ note="chimeric"
483..1142
/ gene="CAT"
483..1142
/ gene="CAT"
/ codon_start=1
/ transl_table=11
/ product="chloramphenicol acetyltransferase"
/ protein_id="AAB39973.1"
/ db_xref="GI:1399736"
/ translation="MEKKITGYTTVDISOWIRKEHFEAFQSVAOCTYNQVOLDITAF
LKTVKKKHKFYPAFIIHLARMAHPELRMAKDGELVIMDSVHPCTVVEHOTETEF
SSLSEYHDDFRQLHYISQDVACYGLENLAFPRKFTENMEFVSANLWVSVFTSFDLNV
ANMNDFFAPVFTMGKYTQGDKVLMLPLAIQVHHAACDGFHVGRLNELQQYCDWEQGG
A"
1173..1394
/ note="SV40 late polyA region"
1414..1650
/ note="SV40"
complement(1708..1727)
/ note="RV primer 4 sequencing primer binding site"
1965
/ note="ColEI-derived plasmid replication origin"
complement(2727..3587)
/ gene="AmpR"
complement(2727..3587)
/ gene="AmpR"
/ function="ampicillin resistance"
/ codon_start=1
/ transl_table=11
/ product="beta-lactamase"
/ protein_id="AAB39974.1"
/ db_xref="GI:1399737"
/ translation="MSIQHFRVALIPFFAAFLPVPFAHPETLVKVKDAEDQLGARVGY
IELDLNSGKILSEFRPEERPMSTFKVLLCGAVLSRIDAGOEQLGRRIRHYQNDLVE
YSPVTEKHLTDGMTVRELCSAAITMSDNTAANLLLTIGGPKELTAFLHMGDHTVRL
DRWPELNEAIPNDERDTPMPVAMATTIRKLITGLTLLASRQQLIDNMEADKVGAPL
LRSLALPAGWFIADKSGAGERSGIITAAALGPDKGPSRIIVITVITGSOATMDERNQIA
EIGASLIKHW"
3720..4174
/ note="fl origin"
4305..4458
/ note="upstream"
4407..4426
/ note="RV primer 3 sequencing primer binding site"
BASE COUNT 1109 a 1101 c 1075 g 1180 t
ORIGIN
Query Match 72.2%; Score 171; DB 12; Length 4465;
Best Local Similarity 100.0%; Pred. No. 5.5e-42;
Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 67 GCTAGCCGGGCTCGAGATCTCGCATCTCAATTAGTCAGCAACCATAGTCCCGC 126
Db 21 GCTAGCCGGGCTCGAGATCTCGCATCTCAATTAGTCAGCAACCATAGTCCCGC 80
Qy 127 CCTAACTCCGCCCATCCGCCCTCACTCGCCAGTTCGCCCATCTCCGCCCATC 186
Db 81 CCTAACTCCGCCCATCCGCCCTCACTCGCCAGTTCGCCCATCTCCGCCCATC 140
Qy 187 GCTGACTAATTTTTTTTATTTATTCAGAGCGCGAGCGCGCTCGGCTCTG 237
Db 141 GCTGACTAATTTTTTTTATTTATTCAGAGCGCGAGCGCGCTCGGCTCTG 191

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LRSLPAGWFIADKSGAGRGSIIAALPGDKPSRIWIYTTGSQATMDERNQIA  
EIGASLIKHW  
misc\_feature 4955..5108  
/note="transcriptional blocker"  
BASE COUNT 1201 a 1392 c 1338 g 1184 t  
ORIGIN

Query Match 72.2%; Score 171; DB 12; Length 5115;  
Best Local Similarity 100.0%; Pred. No. 5.6e-42;  
Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 67 GCTAGCCGGGCTGAGATCTCGGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 126  
|||||  
Db 21 GCTAGCCGGGCTGAGATCTCGGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 80  
|||||  
QY 127 CCCTAACTCCGCCATCCCGCCCTCAACTCCGCCAGTTCGCCCATCTCTCGGCCCATC 186  
|||||  
Db 81 CCCTAACTCCGCCATCCCGCCCTCAACTCCGCCAGTTCGCCCATCTCTCGGCCCATC 140  
|||||  
QY 187 GCTGACTAATTTTTTTATTTATGTCAGAGCGCGAGCGCCCTCGGCCCTCTG 237  
|||||  
Db 141 GCTGACTAATTTTTTTATTTATGTCAGAGCGCGAGCGCCCTCGGCCCTCTG 191  
|||||

RESULT 12  
AX339208  
LOCUS AX339208  
DEFINITION Sequence 2 from Patent WO0196602.  
ACCESSION AX339208  
VERSION AX339208.1 GI:18135469  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.

REFERENCE 1  
AUTHORS Yang, A. L. and Festing, M.  
TITLE Methods and materials to determine the p53 status of a sample by  
determining the binding of p53 to a vector  
JOURNAL Patent: WO 0196602-A 2 20-DEC-2001;  
MEDICAL RESEARCH COUNCIL (GB)  
FEATURES  
source 1..5256  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="pGL3-Control Vector Sequence"

BASE COUNT 1336 a 1268 c 1281 g 1371 t  
ORIGIN

Query Match 72.2%; Score 171; DB 6; Length 5256;  
Best Local Similarity 100.0%; Pred. No. 5.6e-42;  
Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 67 GCTAGCCGGGCTGAGATCTCGGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 126  
|||||  
Db 21 GCTAGCCGGGCTGAGATCTCGGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 80  
|||||  
QY 127 CCCTAACTCCGCCATCCCGCCCTCAACTCCGCCAGTTCGCCCATCTCTCGGCCCATC 186  
|||||  
Db 81 CCCTAACTCCGCCATCCCGCCCTCAACTCCGCCAGTTCGCCCATCTCTCGGCCCATC 140  
|||||  
QY 187 GCTGACTAATTTTTTTATTTATGTCAGAGCGCGAGCGCCCTCGGCCCTCTG 237  
|||||  
Db 141 GCTGACTAATTTTTTTATTTATGTCAGAGCGCGAGCGCCCTCGGCCCTCTG 191  
|||||

RESULT 13  
CVU47296  
LOCUS CVU47296  
DEFINITION Cloning vector pGL3-Control, 5256 bp DNA linear SYN 17-APR-2002  
ACCESSION U47296  
VERSION U47296.2 GI:13195704  
KEYWORDS  
SOURCE Cloning vector pGL3-Control.

## ORGANISM

Cloning vector pGL3-Control  
artificial sequences; vectors.

REFERENCE  
AUTHORS  
TITLE  
JOURNAL

Groskreutz, D.J. and Schenborn, E.T.  
Direct Submission

REFERENCE  
AUTHORS  
TITLE  
JOURNAL

Submitted (26-JAN-1996) D.J. Groskreutz, R&D, Promega Corporation,  
5445 East Cheryl Parkway, Madison, WI 53711, USA

REFERENCE  
AUTHORS  
TITLE  
JOURNAL

Submitted (05-MAR-2001) Technical Writing, Promega Corporation,  
2800 Woods Hollow Road, Madison, WI 53711-5399, USA

REMARK  
COMMENT

Sequence update by submitter  
On Mar 5, 2001 this sequence version replaced gi:1200462.

FEATURES  
source

Location/Qualifiers  
1..5256

/organism="Cloning vector pGL3-Control"

/db\_xref="taxon:45858"

/note="luciferase reporter vector"

misc\_feature

1..41

/note="multiple cloning site"

promoter

48..250

280..1932

/gene="luc"

280..1932

/gene="luc"

/codon\_start=1

/product="luciferase"

/protein\_id="AAA89084.1"

/db\_xref="GI:1200463"

/translation="MEDAKNIKGPAPFPLEDDTAGBQLHKAMRYALVPQTIAFTD  
AHEDVTIAEFEMSVRLAEAMKRYGLNTHRIVVCSENLOFTFMPVLGIFGAV  
AFANDIYNRELLNSGTSQPTVVFVSKGLQILNVQKLPKIIIMDSKDYQV  
FSHARDIFGNQIIPDTALLSVVPHHGFMTTGLYLCGFRVLMVRFERELFLR  
LODYKIQSALLVPTLFSFPAKSTLDKVDLSNLHETASGAPLSKEVEGNAKRHLPL  
GIQGVGLTETTSAILITPEGDKPGKGVVPPFEAKVVDLDTGKTLGVNKGELCV  
RGPIMSGVYNNEPATNALIDKDWLHSGDIAYWDEDEHFFIVDLKSLIKYKGYQVA  
PAELSLILQHENIFDAGVAGLPDDAGELPAAVVLEHGKTMTEKEIVDYVASQVTT  
AKLRGVVVFDEVPKGLTKLDARKIKRELKAKKGGKIAV"

complement(281..303)

/note="GL primer2 sequencing primer binding site"

1964..2185

/note="SV40 late poly(A) signal"

2205..2441

complement(2499..2518)

/note="RV primer4 sequencing primer binding site"

2756

/note="ColEI-derived plasmid replication origin"

complement(3518..4378)

/gene="AmpR"

complement(3518..4378)

/gene="AmpR"

/codon\_start=1

/product="beta-lactamase"

/note="ampicillin resistance"

/protein\_id="AAA89085.1"

/db\_xref="GI:1200464"

/translation="MSIQHFRVALIPFFAFCPLVPAHPETLVKVKDAEDOLGARVGY  
IELDLSNGKILSEFPEERFPMSTFKVLLGAVLSRIDAGAEOLGRIHYSONDLVE  
YSPVTEKHLTDGTVREELCSAAITMSDNTAANLLTTIGGPKELTAFILHNGDHVRL  
DRWPELNEATPNDEPDTMPVAMATTLRLKLTGELLTLASRQQLIDWMEADKRVAGPL  
LRSLPAGWFIADKSGAGRGSIIAALPGDKPSRIWIYTTGSQATMDERNQIA  
EIGASLIKHW"

4510..4965

/note="f1 origin"

5096..5249

/note="upstream poly(A) signal"

5198..5217

/note="RV primer3 sequencing primer binding site"

BASE COUNT 1337 a 1268 c 1281 g 1370 t

ORIGIN

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Query Match      72.2%; Score 171; DB 12; Length 5256;
Best Local Similarity 100.0%; Pred. No. 5.6e-42;
Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 67 GCTAGCCGGCTCGAGATCTGCATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 126
    |||
DB 21 GCTAGCCGGCTCGAGATCTGCATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 80

QY 127 CCCTAACTCCGCCATCCCGCCCTAACTCCGCCAGTTCGCCGCCATCTCTCGGCCCATC 186
    |||
DB 81 CCCTAACTCCGCCATCCCGCCCTAACTCCGCCAGTTCGCCGCCATCTCTCGGCCCATC 140

QY 187 GCTGACTAATTTTTTTTATTTATGACAGAGCCGAGCGCCCTCGGCTCTG 237
    |||
DB 141 GCTGACTAATTTTTTTTATTTATGACAGAGCCGAGCGCCCTCGGCTCTG 191

RESULT 14
AB038600
LOCUS      AB038600      6320 bp      DNA      circular SYN 22-FEB-2001
DEFINITION Cloning vector pVLUC441 DNA, complete sequence.
ACCESSION  AB038600
VERSION    AB038600.1 GI:13094136
KEYWORDS  beta-lactamase; luciferase.
SOURCE     Cloning vector pVLUC441 DNA.
ORGANISM   artificial sequences: vectors.
REFERENCE  1 (sites)
AUTHORS    Hashinaka,K.
TITLE      Synthetic Autonomous Vectors Based on Palindromic Sequences of
           Parvovirus B19
JOURNAL    Published Only in DataBase (2001)
REFERENCE  2 (bases 1 to 6320)
AUTHORS    Hashinaka,K.
TITLE      Direct Submission
JOURNAL    Submitted (21-FEB-2000) Kazuya Hashinaka, Miyazaki Medical College,
           Department of Biochemistry; 5200 Kihara, Kiyotake, Miyazaki
           889-1692, Japan (E-mail:hashinaka@postl.miyazaki-med.ac.jp,
           Tel:81-985-85-0985, Fax:81-985-85-2401)

FEATURES             Location/Qualifiers
     source          1..6320
                     /organism="Cloning vector pVLUC441"
                     /db_xref="taxon:117918"
                     /focus
     source          6..479
                     /organism="B19 virus"
                     /note="xref=taxon:10798"
                     /notes="synonym:Parvovirus B19"
     source          2937..3420
                     /organism="B19 virus"
                     /note="xref=taxon:10798"
                     /notes="synonym:Parvovirus B19"
     repeat_region   9..391
     gene            1347..2414
                     /gene="luc"
                     /gene="luc"
                     /codon_start=1
                     /transl_table=11
                     /product="luciferase"
                     /protein_id="BAB32737.1"
                     /db_xref="GI:13094137"
                     /translation="MNSGSGTGLPKGVLPHTACVRFSHARDPIFGNIIPTDAIL
                     VPFHFGFTGLGICFRVLMYREAEFLRLSDYKIQSALIVPTLFSFPAK
                     STLDYDLSNLHIAAGGAPLSKEVEAKRFLPGROGYLTETTSAILTPEAG
                     DDPAGVKVPFFFAKVLDLTGKTLGYNQRCGLCVRGPMIMSGYVNNPEATNALID
                     KDWLHSGDIAYDEDEHEFFIVDLRLSLIKYKIQVAPAELESILLQHPNIFDAGVAC
                     LPDDAGCELPAAVVLEHGKTMTEKEIVDYVASQVTTAKKRGGVVFDVEPKGLTGK
                     LDARKIREILIKAKGKTIAY"
     CDS              3029..3411
                     /codon_start=1
                     /transl_table=11
                     /gene="ampR"
                     /product="ampR"
                     /product="modified C"

Query Match      67.5%; Score 160; DB 12; Length 6320;
Best Local Similarity 99.4%; Pred. No. 1.4e-38;
Matches 171; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 67 GCTAGCCGGCTCGAGATCTGCATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 126
    |||
DB 495 GCTAGCCGGCTCGAGATCTGCATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGC 554

QY 127 CCCTAACTCCGCCATCCCGCCCTAACTCCGCCAGTTCGCCGCCATCTCTCGGCCCAT 185
    |||
DB 555 CCCTAACTCCGCCATCCCGCCCTAACTCCGCCAGTTCGCCGCCATCTCTCGGCCCAT 614

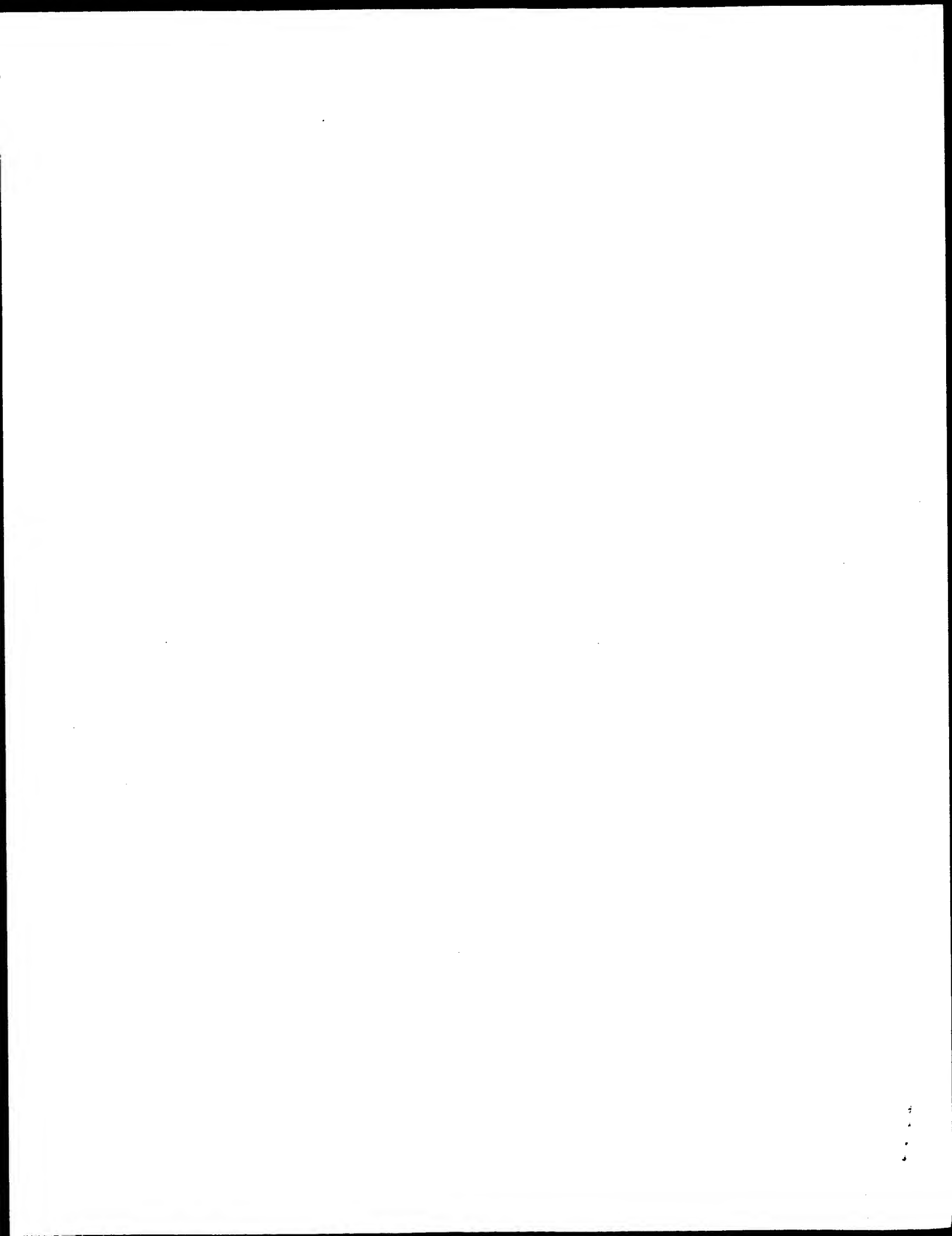
QY 186 GCCTGACTAATTTTTTTTATTTATGACAGAGCCGAGCGCCCTCGGCTCTG 237
    |||
DB 615 GCCTGACTAATTTTTTTTATTTATGACAGAGCCGAGCGCCCTCGGCTCTG 666

RESULT 15
AB037684
LOCUS      AB037684      5917 bp      DNA      circular SYN 04-APR-2000
DEFINITION Cloning vector pPLuc, complete sequence.
ACCESSION  AB037684
VERSION    AB037684.1 GI:7415874
KEYWORDS  Pluc; luciferase; modified C.
SOURCE     synthetic construct DNA.
ORGANISM   artificial sequences.
REFERENCE  1 (bases 1 to 5917)
AUTHORS    Chen,A.B., Kao,A.Y.-F. and Brown,C.M.
TITLE      A short open reading frame within the encapsidation signal affects
           the translation of the polymerase gene from hepatitis Bvirus
JOURNAL    Unpublished
REFERENCE  2 (bases 1 to 5917)
AUTHORS    Chen,A.B., Kao,A.Y.-F. and Brown,C.M.
TITLE      Direct Submission
JOURNAL    Submitted (27-JAN-2000) Chris M. Brown, University of Otago,
           Department of Biochemistry; P.O.Box 56, Dunedin, Otago 9001, New
           Zealand (E-mail:chris.brown@stonebow.otago.ac.nz,
           Tel:+64-3-479-7875, Fax:+64-3-479-7866)

FEATURES             Location/Qualifiers
     source          1..5917
                     /organism="synthetic construct"
                     /db_xref="taxon:32630"
                     /note="cloning vector pPLuc"
     misc_feature    1..495
                     /note="HBV Adw 5' leader of pregenomic RNA"
     CDS              35..94
                     /note="ORF"
                     /codon_start=1
                     /evidence=experimental
                     /transl_table=11
                     /protein_id="BAA93573.1"
                     /db_xref="GI:7415875"
                     /translation="MSHCSSLQAVPVALGHGH"
     CDS              84..581
                     /codon_start=1
                     /transl_table=11
                     /product="modified C"

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GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: January 3, 2003, 22:54:17 ; Search time 3151.28 Seconds  
(without alignments)  
1218.024 Million cell updates/sec

Title: US-09-787-562-9

Perfect score: 237  
Sequence: I gctagagctctgcaggacgt.....cgaggcgcctcgccctctg 237

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

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1: em_estba:*
2: em_esthum:*
3: em_estlin:*
4: em_estmu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_esthc:*
9: gb_estli:*
10: gb_est2:*
11: gb_hlc:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_inv:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_mam:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rod:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	80.8	34.1	569	12 BG063359	BG063359 H3006C07-
C 2	79.8	33.7	392	9 AA078277	AA078277 7P01B01 C
C 3	55	23.2	578	9 AA409312	AA409312 EST03748
C 4	41	17.3	869	10 BE216820	BE216820 HV_CEB001
C 5	38.2	16.1	974	17 A2157908	A2157908 SP_0059_A
C 6	37.6	15.9	868	17 CNS03VAE	AL262175 Tetraodon

C	7	37.2	15.7	684	12	BG620549
	8	37.2	15.7	725	17	AG062985
	9	37	15.6	639	17	B19583
C	10	37	15.6	862	17	AG137051
	11	37	15.6	1619	14	BM906235
	12	36.8	15.5	371	13	B1885730
	13	36.8	15.5	911	17	AZ682313
C	14	36.4	15.4	522	17	CNS015B2
C	15	36.4	15.4	978	17	CNS018FK
	16	36.2	15.3	453	13	B1885726
	17	36.2	15.3	518	17	AZ079405
	18	36.2	15.3	830	17	B19180
	19	36.2	15.3	1101	17	CNS0181N
	20	36	15.2	552	17	B19569
C	21	36	15.2	629	10	BE239175
C	22	36	15.2	631	10	AW727988
	23	36	15.2	982	17	CNS014D5
	24	36	15.2	1137	13	BM415579
	25	35.6	15.0	755	17	AG100419
C	26	35.6	15.0	1087	17	AG147408
	27	35.6	15.0	1101	17	CNS0036H
	28	35.4	14.9	309	13	B1885645
	29	35.4	14.9	866	12	BG442104
	30	35.2	14.9	625	17	B19473
	31	35.2	14.9	870	17	BH157873
	32	35.2	14.9	892	17	BH133618
C	33	35.2	14.9	929	17	AZ687353
	34	35.2	14.9	971	17	BH156534
	35	35.2	14.9	1093	13	BM544531
	36	35	14.8	398	12	BG364630
	37	35	14.8	449	13	B1885612
C	38	35	14.8	575	9	AA800583
	39	35	14.8	802	17	AG139568
	40	35	14.8	846	17	B18514
	41	35	14.8	954	13	BF833711
	42	35	14.8	1192	17	B11834
	43	35	14.8	1211	14	BM807956
	44	35	14.8	1389	14	BM924250
	45	34.8	14.7	657	17	AG111841

#### ALIGNMENTS

RESULT 1 BG063359/c  
LOCUS H3006C07-3 NTA Mouse 15K cDNA Clone Set Mus musculus cDNA clone  
DEFINITION H3006C07 3', mRNA sequence.  
ACCESSION BG063359  
VERSION BG063359.1 GI:12545922  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.  
AUTHORS 1 (bases 1 to 569)  
T.S., Carter, M.G. and Ko, M.S.H.  
TITLE Verification and initial annotation of NIA mouse 15K cDNA clone set  
JOURNAL Unpublished (2001)  
COMMENT Contact: George J. Kargul  
Laboratory of Genetics  
National Institute on Aging/National Institutes of Health  
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA  
Email: cdaa@igsun.grc.nia.nih.gov  
This clone set has been freely distributed to the community. Please visit <http://igsun.grc.nia.nih.gov/cDNA/15K.html> for details.  
Plate: H3006 row: C column: 07  
Seq primer: -21M13 Forward  
High quality sequence stop: 569  
POLYA-No.

#### FEATURES

Location/Qualifiers

2

uninoculated leaves were harvested 20 hr post-inoculation (Wei, Wise). In the TJ Close lab at the University of California, Riverside, total RNA was prepared from each sample pool, equal quantities of all three RNA pools were combined, poly(A) RNA was purified from the mixture, one primary unamplified cDNA library was made, and 1 million pfu were in vivo excised to give pBluescript SK(-) cDNA phagemids (Choi, Close). Phagemids were plated and picked at the Clemson University Genomics Institute (CUGI) (Begum , Palmer, Frisch, Atkins and Wing). Plasmid DNA preparations, DNA sequencing and sequence analysis were performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates , Rambo, Main). The sequence has been trimmed to remove vector sequence and contains a minimum of 100 bases of phred value 20 or above. For more details on library preparation and sequence analysis see <http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> Also see Clouse TJ, Wang R, Kleinhofs A, Wise R (2001) Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html>)"

BASE COUNT      174 a    213 c    367 g    115 t

ORIGIN

Query Match                          17.3%; Score 41; DB 10; Length 869;  
Best Local Similarity       58.7%; Pred. No. 3.3;  
Matches       71; Conservative       0; Mismatches       50; Indels       0; Gaps       0;

QY    115 CCATAGTCCGCCCTTAACTCGCCCATCGCCGCCCCCTAATCGCCGCAGTTCCGGCCATT 174  
|| | ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||  
Db    837 CCTTCCGCCGCCGCCCTACCCTCCGCCCTCCGCCCTCCGCCCTTTCCCTCCTCT 778  
  
QY    175 CTCGGCCCCATCGCTGACTAAATTTTTTTATTTATCGAGAGCGGAGCGCCTCGGCCCT 234  
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||  
Db    777 CTCTCCCTCCCGCACTACCCCTTATTACTTTTCACCGCGCCCTCGCCCACCTCC 718  
  
QY    235 C 235  
||  
Db    717 C 717

RESULT 5

AZ157908

LOCUS

DEFINITION

SP\_0059.A1.D06.T7A Strongylocentrotus purpuratus, purple sea urchin sperm genomic BAC library Strongylocentrotus purpuratus genomic clone plate=59 Col=11 Row=G, DNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

GSS.  
AZ157908.1 GI:8310511  
Strongylocentrotus purpuratus.  
Strongylocentrotus purpuratus  
Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa; Echinoidea; Euechinoidea; Echinacea; Echinoida;  
Strongylocentrotidae; Strongylocentrotus.  
1 (bases 1 to 974)  
Cameron,R.A., Mahairas,G., Rast,J.P., Martinez,P., Biondi,T.R., Swartzell,S., Wallace,J.C., Poustka,A.J., Livingston,B.T., Wray,G.A., Ettensohn,C.A., Lehrach,H., Britten,R.J., Davidson,E.H. and Hood,L.  
A sea urchin genome project: Sequence scan, virtual map, and additional resources  
Proc. Natl. Acad. Sci. U.S.A. 97 (17), 9514-9518 (2000)

TITLE

JOURNAL

MEDLINE

COMMENT

Contact: Cameron, RA, Davidson, EH, Hood, L  
Division of Biology 156-29  
California Institute of Technology  
Pasadena California 91125, USA  
Tel: (626) 395-8421  
Fax: (626) 793-3047  
Email: acameron@caltech.edu  
Plate: 59 row: G column: 11

BASE COUNT	188 a	187 c	234 g	232 t	27 others
ORIGIN					

Query Match	15.9%	Score 37.6;	DB 17;	Length 868;
Best Local Similarity	55.4%	Pred. No. 23;		
Matches 62; Conservative		5; Mismatches	45; Indels	0; Gaps

QY 111 GCAACCATAGTCCCGGCCCTAACTCGGCCCATCCCGGCCCTAACTCGGCCAGTTCGGCC 170

171 CATTCTCCGCCCATCGCTGACTAATTTTTTTTATTTATGCAGAGGCCGAGG 222

Db  
576 TCDTCCTCTCCCAAAHTTATWTATWAAWAAAGTTAGGAAGCGGAGGTGG 525

RESULT	7
BG620549/c	
LOCUS	684 bp mRNA linear EST 18-APR-2001
DEFINITION	603619703f1 NIH_MGC_79 Homo sapiens CDNA IMAGE:473280 5', mRNA sequence.

ACCESSION	BG620549	GI:13671920
VERSION	BG620549.1	
KEYWORDS	EST.	
SOURCE	human.	

ORGANISM      Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE	1 (bases 1 to 684)
AUTHORS	NIH-MGC <a href="http://mgc.uci.nih.gov/">http://mgc.uci.nih.gov/</a>
TITLE	National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL	Unpublished (1999)
COMMENT	Contact: Robert Strausberg Ph.D.

Email: [cgapbs-r@mail.nih.gov](mailto:cgapbs-r@mail.nih.gov)  
Tissue Procurement: CLONTECH Laboratories, Inc.  
DNA Library Preparation: CLONTECH Laboratories, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
Plate: LLCM1594 row: j column: 01  
High quality sequence stop: 404.

FEATURES	Source
Location/Qualifiers	
1. 684	
/organism="Homo sapiens"	
/db_xref="taxon:9606"	
/clone_image="733280"	
/clone_lib="NIH_MGC_79"	
/lab_host="DH10B (T1 phage-resistant)"	
/note="Organ: placenta; Vector: pDNR-LIB (Clontech); Site:1: SfiI (ggcccttcggc); Site:2: SfiI (ggccattatgcc) ; 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGGCATTATGCC-3' and 3' adaptor sequence: 5'-ATCTAGAGCGGAGCGGCACATG-dT(30)BN-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.3 kb (range 0.5-4.0 kb). 15/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA). Note: this is a NIH_MGC Library."	

BASE COUNT	258 a	160 c	207 g	59 t
ORIGIN				
Query Match		15.7%	Score 37.2;	DB 12; Length 684;
Best Local Similarity		54.3%	Pred. No. 28;	
Matches 75; Conservative		0;	Mismatches 63;	Indels 0; Gaps

[illegible]

Seq primer: T7  
Class: BAC ends  
High quality sequence stop: 974.

FEATURES

source	Location/Qualifiers
1..974	

```

/organism="Strongylocentrotus purpuratus"
/db_xref="taxon:7668"
/clone="plate=59 Col=11 Row=G"
/clone_lib="Strongylocentrotus purpuratus, purple sea
urchin, sperm genomic BAC library"
/note="Organ: sperm; Vector: BACE3.6; BAC Clones in E-Coli
DH10B"

```

BASE COUNT	240 a	413 c	123 g	198 t
ORIGIN				

Query Match	16.1%;	Score 38.2;	DB 17;	Length 9/4;
Best Local Similarity	58.3%;	Pred. No. 16;		
Watchdog 57. Conservative		0: Mismatches	48:	Indels
		0:	0:	Gaps
				0:

[illegible]

132 ACTCGGCCCATCCCGCCCCCTAACTCGGCCAGTTCCGCCCATTTCTCGGCCCATC 186

Db  
886 CCCCCCACACCTTCCCCATACTCCCCCATCTCCCCCCCCCCCCACCCCCACC 940

RESULT 6	
CNS03VAE/c	CNS03VAE
LOCUS	868 bp DNA linear GSS 18-MAY-2000
DEFINITION	Tetraodon nigroviridis genome survey sequence PUC-Orig of clone 061P09 of library G from Tetraodon nigroviridis, genomic survey sequence.

AL262175  
AL262175.1  
GI:7983801  
GSS: genome survey sequence.  
Tetradodon nigroviridis.  
SOURCE

ORGANISM

Tetraodon nigroviridis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;  
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;  
Tetraodontidae; Tetraodon.

1 (bases 1 to 888)  
Roest-Crolius H., Jaillon O., Dasilva C., Bouneau L., Fisher C.,  
Bernot A., Fitzames C., Wincker P., Brottier P., Quetier F.,  
Saurin W. and Weissenbach J.  
Human gene number estimate provided by genome wide analysis using  
mouse and human orthologous DNA sequence

2 (bases 1 to 868)  
Roest-Crollius, H., Jaillon, O., Dasilva, C., Fizames, C., Fisher, C.,  
Bonneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and  
Weissenbach, J.

TITLE	Characterization and repeat analysis of the compact genome of the freshwater pufferfish <i>Tetraodon nigroviridis</i>
JOURNAL	Unpublished
REFERENCE	3 (bases 1 to 868)
AUTHORS	Genoscope.

TITLE	JOURNAL	COMMENT
Direct submission		
Submitted (12-APR-2000)		
This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetraodon nigroviridis genome. For more information, please take a look at <a href="http://www.genoscope.cns.fr/tetraodon">http://www.genoscope.cns.fr/tetraodon</a>		

```

FEATURES
source
1 968
Location/Qualifiers
/organism="Tetraodon nigroviridis"
/ab_xref="taxon:99883"
/clone="061809"
/clone_lib="C"
/note="Genoscope sequence ID : COBG061CH05SP1-end :

```



```

Db 490 GCGGCCCCCCCCCGGCTTCGGCGCGCTCGGCGCGCGCGCGCGCGCGCGCGCTCC 431
QY 192 CTAATTTTTTTTATTTAT 209
Db 430 TGCTTTTTTTTCTTCT 413

RESULT 8
AG062985
LOCUS Pan troglodytes DNA, clone: PTB-051101.F, genomic survey sequence.
DEFINITION AG062985
ACCESSION AG062985
VERSION AG062985.1 GI:16614787
KEYWORDS GSS.
SOURCE Pan troglodytes male lymphoblast DNA, clone_lib:PTB Chimpanzee Male
BAC Library clone:PTB-051101.F.
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
REFERENCE
AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE BAC end sequences of Library PTB
JOURNAL Unpublished
REFERENCE
AUTHORS 2 (bases 1 to 725)
Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE Direct Submission
JOURNAL Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suehiro-chou, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail: chimbes@gsc.riken.go.jp, URL: http://hgp.gsc.riken.go.jp/,
Tel: 81-45-503-9111, Fax: 81-45-503-9170)
COMMENT Clones are derived from the chimpanzee BAC library PTB This BAC end
was generated during the R&D process and may have higher chance of
clone tracking errors.
PRIMERS
Sequencing: -21M13
LIBRARY
Vector : pKSL45
R.Site 1 : SacI
R.Site 2 : SacI.
FEATURES
source
Location/Qualifiers
1..725
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/clone="PTB-051101.F"
/sex="male"
/cell_type="lymphoblast"
/clone_lib="PTB Chimpanzee Male BAC Library"
BASE COUNT 55 a 423 c 51 g 195 t 1 others
ORIGIN

Query Match 15.7%; Score 37.2; DB 17; Length 725;
Best Local Similarity 57.9%; Pred. No. 28;
Matches 66; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 92 TCTGCATCTCAATTAGTCAGCAACCATAGTCCGCGCCCTTAACCTCGGCCCATCCGCGCCCT 151
Db 550 TCCTTCCTCCCTCTTCCTCCGCGCCCTCTCTCCCTCCGCGCCCTCTCTCCCTCCGCGCCCT 609
QY 152 AACCTCCGCGCGAGTTCGCGCCCATCTCGCGCCCATCGCTGACTTAATTTTTTTAT 205
Db 610 TCCTTTTCCCGCTCCCGCTCTCTCCGCGCCCTCTCTCTCTCTCTCTCTCTCTCTCTCT 663

RESULT 9
B19583
LOCUS Pan troglodytes DNA, clone: PTB-150N02.F, genomic survey sequence.
DEFINITION B19583
ACCESSION B19583
VERSION B19583.1 GI:16666729
KEYWORDS GSS.
SOURCE Pan troglodytes male lymphoblast DNA, clone_lib:PTB Chimpanzee Male
BAC Library clone:PTB-150N02.F.
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
REFERENCE
AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE BAC end sequences of Library PTB
JOURNAL Unpublished

```

```

B19583.1 GI:2394637
GSS.
thale cress.
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 639)
Feng, J., Dewar, K., Buehler, E., Kim, C., Li, Y., Shinn, P., Sun, H. and
Ecker, J.
BAC End Sequences at ATGC
Unpublished (1997)
Other GSSs: F1114-Sp6.1, F1114-Sp6, F1114-Sp6.2, F1114-Sp6.3,
F1114-Sp6.4
Contact: Ecker, J.
Arabidopsis Thaliana Genome Center
University of Pennsylvania
Dept. of Biology, University of Pennsylvania, Philadelphia, PA
19104
Tel: 215-898-9384
Fax: 215-898-8780
Email: jecker@atgenome.bio.upenn.edu
Seq primer: T7
Class: BAC ends
High quality sequence start: 458
High quality sequence stop: 476.
Location/Qualifiers
1..639
/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="F1114"
/clone_lib="IGF"
/sex="hermaphrodite"
/note="Vector: BelobAC11; Site_1: EcoRI; Site_2: EcoRI;
Produced by Thomas Altmann"
BASE COUNT 81 a 365 c 32 g 130 t 31 others
ORIGIN

Query Match 15.6%; Score 37; DB 17; Length 639;
Best Local Similarity 54.7%; Pred. No. 32;
Matches 70; Conservative 0; Mismatches 58; Indels 0; Gaps 0;

QY 108 TCAGCAACCATAGTCCGCGCCCTAAGTCCGCGCCATCCGCGCCCTACTCGGCCAGTTCC 167
Db 373 TCTCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCC 432
QY 168 GCCATTCCTCCGCGCCCATCGCTGACTTAATTTTTTTATTTATGAGAGCCGAGCGCCGC 227
Db 433 NCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCCCTCC 492
QY 228 TCGGCTC 235
Db 493 CCNCCCTCC 500

RESULT 10
AG137051/c
LOCUS Pan troglodytes DNA, clone: PTB-150N02.F, genomic survey sequence.
DEFINITION AG137051
ACCESSION AG137051
VERSION AG137051.1 GI:16666729
KEYWORDS GSS.
SOURCE Pan troglodytes male lymphoblast DNA, clone_lib:PTB Chimpanzee Male
BAC Library clone:PTB-150N02.F.
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
REFERENCE
AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE BAC end sequences of Library PTB
JOURNAL Unpublished

```

[illegible]

```

/organism="Danio rerio"
/db_xref="taxon:7955"
/clone_lib="4490198"
/clone_lib="zebrafish Washu MPING EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield
stage embryos"
/lab_host="X11-blue MRF"
/Note="Vector: pSPORT1; Site_1: NotI; Site_2: SalI; 1st
strand cDNA was primed with a Not I - oligo(dT)15 primer
[5'-GCAGTACTTCTAGATCGGAGCGCCGCTTTTCTTTT3'];
double-stranded cDNA was ligated to Sal I adaptors (BRL),
digested with Not I and cloned into the Not I and Sal I
sites of the pSPORT1 vector (BRL). Library was constructed
by Matthew Clark (Lehrach lab; ICRF, London and Max Planck
Institut fuer Molekulare Genetik, Berlin). cDNAs for EST
analysis were selected following oligonucleotide
hybridization fingerprinting of arrayed clones from
zebrafish late somitogenesis (26 ss), adult liver or
embryonic shield stage (5.6 h) libraries. Fingerprint
data were used to computationally cluster cDNAs, and a
single cDNA from each cluster was chosen for sequencing.
In some cases multiple members of the same cluster were
sequenced to assess clustering parameters or single clones
were sequenced additional times to assess quality
control."
BASE COUNT      5 a 298 c 6 g 37 t 25 others
ORIGIN

Query Match      15.5%; Score 36.8; DB 13; Length 371;
Best Local Similarity 57.0%; Pred. NO. 34;
Matches 65; Conservative 0; Mismatches 49; Indels 0; Gaps 0;

QY 122 CCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCC 181
Db 128 CCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCC 187

QY 182 CCATCGCTGACTAATTTTATTTATTTATTTATTTATTTATTTATTTATTTATTTATTTAT 235
Db 188 CCGCGCCCGCTTTTATTTTATTTATTTATTTATTTATTTATTTATTTATTTATTTATTTAT 241

RESULT 13
A2682313
LOCUS
DEFINITION
  ENFKQ34TF Entamoeba histolytica Sheared DNA linear GSS 14-DEC-2000
  genomic, DNA sequence.
ACCESSION
  A2682313
KEYWORDS
  GSS.
SOURCE
  Entamoeba histolytica.
  Entamoebidae; Entamoeba.
REFERENCE
  1 (bases 1 to 911)
  Loftus,B., Van Aken,S. and Fraser,C.
  Determination of clone end sequences from Entamoeba histolytica
  HMI:IMSS sheared DNA library
  Unpublished (2000)
  Contact: Brendan J Loftus
  Department of Eukaryotic Genomics
  The Institute for Genomic Research
  9712 Medical Center Dr., Rockville, MD 20850, USA
  Tel: 301 838 0208
  Fax: 301 838 3543
  Email: bjloftus@tigr.org
  Clones are derived from the Entamoeba histolytica HMI:IMSS sheared
  DNA library
  Seq primer: M13-Forward
  Class: shotgun
  High quality sequence start: 30
  High quality sequence stop: 330.
  High quality sequence
  Location/Qualifiers
    1..911

```

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/organism="Entamoeba histolytica"
/strain="HMI:IMSS"
/db_xref="taxon:5759"
/clone_lib="Entamoeba histolytica Sheared DNA"
/Note="Vector: PHOSI; Site_1: Bst I; Constructed at The
Institute for Genomic Research (TIGR), Rockville, MD.
Genomic DNA isolated from broth cultures of E. histolytica
using a method described by Clark and Diamond (Clark,
C.G., and Diamond, L.S. (1993) Entamoeba histolytica: a
method for isolate identification. Exp. Parasitol.
77:450.). The DNA was mechanically sheared to give a
tight size distribution (~2 kb). The v + i method used for
the library construction is described in detail in Smith,
H.O. and Venter, J.C. (Making small insert libraries for
whole genome shotgun sequencing projects. In Genome
Sequencing: A Practical Approach, eds. M. Vaudin and B.
Barell, Oxford University Press, 1999)."
BASE COUNT      171 a 117 c 74 g 549 t
ORIGIN

Query Match      15.5%; Score 36.8; DB 17; Length 911;
Best Local Similarity 63.6%; Pred. No. 36;
Matches 56; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

QY 122 CCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCCCTAAGTCCGCGCC 181
Db 373 CCGCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCC 432

QY 182 CCATCGCTGACTAATTTTATTTATTTATTTATTTATTTATTTATTTATTTATTTATTTAT 209
Db 433 CCGCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCC 460

RESULT 14
CNS015BZ/c
LOCUS
DEFINITION
  Drosophila melanogaster genome survey sequence sp6 end of BAC
  BACN13C14 of DrosBAC library from Drosophila melanogaster (fruit
  fly), genomic survey sequence.
ACCESSION
  AL105257
VERSION
  AL105257.1 GI:5617271
KEYWORDS
  GSS.
SOURCE
  Drosophila melanogaster.
  Drosophila melanogaster.
  Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
  Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
  Ephydroidea; Drosophilidae; Drosophila.
REFERENCE
  1 (bases 1 to 522)
  Genoscope.
  Direct Submission
  Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :
  BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
  - Web : www.genoscope.cns.fr)
  Determination of this BAC-end sequence was carried out as part of a
  collaboration with the European Drosophila Genome Project (EDGP) -
  http://www.edgp.ebi.ac.uk. This Drosophila melanogaster BAC
  library (Dros BAC) was made by Alain Billaud at CPH (Centre
  d'Etude du Polymorphisme Humain) with funding provided by a MRC
  project grant. The DNA was prepared from embryos by Alain Bucheton
  and Genevieve Payan. It has been constructed in the vector
  pBelobAC11.
FEATURES
  Location/Qualifiers
    1..522
    /organism="Drosophila melanogaster"
    /db_xref="taxon:7227"
    /clone_lib="BACN13C14"
    /clone_lib="DrosBAC"
    /plasmid="pBelobAC11"
    /note="end : SP6"
BASE COUNT      89 a 32 c 303 g 47 t 51 others
ORIGIN

Query Match      15.4%; Score 36.4; DB 17; Length 522;

```

Search completed: January 4, 2003, 01:04:18  
Job time : 3158.28 secs

Result No.	Score	Query %		Length	DB	ID	Description
		Match					
1	237	100.0	237	21	AA12001		Murine PGK HRE der
2	221	93.2	243	20	AA211397		Murine PGK fragmen
3	221	93.2	243	21	AA11995		Murine PGK HRE der
4	209	88.2	242	20	AA207789		Promoter OBhrel us
5	175	73.8	242	21	AA12016		Murine PGK HRE der
6	171	72.2	5010	24	AA227539		pGL3 promoter vect
7	171	72.2	5256	21	AA207776		DNA sequence of pl
8	171	72.2	5256	24	AA227537		pGL3 control vecto
9	152.2	64.2	267	21	AA12019		Human erythropoiet

Novel polynucleotide constructs comprising at least two repeats of a hypoxia response element useful for driving expression of nucleic acids of interest in a cell under hypoxic conditions

Example 1; Page 68; 155pp; English.

This invention describes novel polynucleotide comprising at least 2 repeats of a hypoxia response element (HRE), where the hypoxia-inducible factor (HIF) consensus binding sites within each of the 2 repeats are separated by a spacer of at least 20 contiguous nucleotides. The products of the invention have vasotropic, cardiatic, cytostatic and antiarthritic activity and can be used for gene therapy. The polynucleotides are useful for delivering nucleic acids of interest to mammalian cells. Lentiviral vectors are responsive to hypoxic agents and to agents that mimic hypoxia. This regulation can be harnessed in vitro to enhance the production of the vector and can be used in vivo to regulate gene expression in response to a physiological signal. The vectors have utility in response to hypoxia, where ischaemia, including hypoxia, is a feature, e.g. cardiovascular disease, peripheral arterial disease, cancer and arthritis. The novel regulatory construct is capable of driving very high levels of transcription under conditions of hypoxia whilst providing only low basal levels of transcription under normal oxygen conditions. The polynucleotide construct targets cells within a tumor mass that are under conditions of hypoxia without affecting normal surrounding tissue. This sequence represents a murine phosphoglycerate kinase (PGK) HRE derived construct which is described in the method of the invention.

Query Match	100.0%;	Score 237;	DB 21;	Length 237;
Best Local Similarity	100.0%;	Pred. No. 6.3e-63;		
Mismatches 237: Conservative	0;	Mismatches	0;	Indels 0;
Gaps	0;			

1	GCTAGAGTCGTGCAGACGTGCATCTAGTGTCTGTCGAGCATCTAGTGTCTGTCGAGGAC	60
QY		
1	GCTAGAGTCGTGCAGACGTGCATCTAGTGTCTGTCGAGGACATCTAGTGTCTGTCGAGGAC	60
Db		
61	CTGACAGCTAGCCCGGCTCGAGATCTCGGATCTGCATCTCAATTAGTCAGCAACCATAG	120
QY		
61	GTGACAGCTAGCCCGGCTCGAGATCTGGATCTGCATCTCAATTAGTCAGCAACCATAG	120
Db		
121	TCCGGCCCTAACTTCGCGCCCATCCCGCCCTAACTTCGCGCCAGTTCGCGCCATCTTCGCG	180
QY		
121	TCCGGCCCTAACTTCGCGCCCATCCCGCCCTAACTTCGCGCCAGTTCGCGCCATCTTCGCG	180
Db		
181	CCCAATCGTGACTAAATTTTTTTTATTCGAGAGCGCGCGCTCGGCTCTG	237
QY		
181	CCCAATCGTGACTAAATTTTTTTTATTCGAGAGCGCGCGCTCGGCTCTG	237
Db		

RESULT	2
AAZ11397	
ID	AAZ11397 standard; DNA; 243 BP.
XX	
XX	
AC	AAZ11397;
XX	
XX	
DT	26-OCT-1999 (first entry)
XX	
XX	
DE	Murine pCk fragment linked to SV40 promoter.

23-SEP-1998; 98NO-GB02885.

25-SEP-1997; 97GB-0020465.

23-SEP-1997; 97GB-0020216.

(OXFO-) OXFORD BIOMEDICA UK LTD.

Bebbington C, Binley KM, Lewis C, Naylor S;  
WPI; 1998-263482/22.

New retroviral vectors, for, e.g. delivering nucleotide sequences to solid tumor sites

Example 1; Page 69; 288pp; English.

The invention relates to a retroviral vector (RV) comprising a functional splice donor site (FSDS) and a functional splice acceptor site (FSAS) where: (i) the FSDS and the FSAS flank a first nucleotide sequence of interest (NOI); (ii) the FSDS is upstream of the FSAS; (iii) the RV is derived from a retroviral pro-vector; (iv) the retroviral pro-vector comprises a first nucleotide sequence (NS) capable of yielding the FSDS and a second NS capable of yielding the FSAS; and (v) the first NS is downstream of the second NS, such that the RV is formed as a result of reverse transcription of the retroviral pro-vector. A hybrid viral vector (VV) system for in vivo gene delivery, which system comprises a primary VV which encodes a secondary VV, the primary vector capable of infecting a first target cell and of expressing the secondary VV, which secondary vector is capable of transducing a secondary target cell, where the primary vector is obtainable from or is based on an adenoviral vector and the secondary VV is obtainable from or is based on a RV preferably a lentiviral vector (LVV) is also provided. The systems can be used for delivering NOIs to one or more target sites. The NOIs may encode therapeutic or diagnostic agents. The methods are used particularly for producing modified hematopoietic stem cells (MHSCs) to deliver NOIs to sites such as solid tumors which are characterised by ischemia, such as hypoxia or low glucose concentration. The system permits the stable expression of NOIs in targeted cells, e.g. rapidly dividing cells. The present sequence represents a trimer encompassing -307/-290 sequence of murine PGK, linked to SV40 promoter. The sequence comprises murine PGK hypoxia response element (HRE) sequence.

```

Query Match      93.2%; Score 221; DB 20; Length 243;
Best Local Similarity 97.5%;
Pred. No. 5e-58;
Matches 237; Conservative 0; Mismatches 0; Indels 6; Gaps 1;

```

QY	1	GCTAGAGTCGTGCAGGACGTGACATCTAGTGTGTCAGG-----CATCTAGTGTCTGTG	54
Db	1	GCTAGAGTCGTGCAGGACGTGACATCTAGTGTGTCAGGACGTGACATCTAGTGTCTGTG	60
QY	55	CAGGAGCTCACAGCTAGCCCGGCTCGAGATCTGCATCTCAATTAGTCAGCAA	114
Db	61	CAGGAGCTGACAGCTAGCCCGGCTCGAGATCTGCATCTCAATTAGTCAGCAA	120
QY	115	CCATAGTCCGGCCCTTAATCTCGCCCATCCCGGCCCTTAATCTCCGCCAGTTCGCCGCATT	174
Db	121	CCATAGTCCGGCCCTTAATCTCGCCCATCCCGGCCCTTAATCTCCGCCAGTTCGCCGCATT	180
QY	175	CTCCGCCCATCGCTGACTAATTTTTTTTATTATGTCAGAGCCGAGGCCGCTCGGCT	234
Db	181	CTCCGCCCATCGCTGACTAATTTTTTTTATTATGTCAGAGCCGAGGCCGCTCGGCT	240
QY	235	CTG	237
Db	241	CTG	243

RESULT 3  
AAA11995  
ID AAA11995 standard; DNA; 243 BP.

```

XX AC AAA11995;
XX DT 14-AUG-2000 (first entry)
XX DE Murine PGK HRE derived promoter OBHrel DNA.
XX KW HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;
XX KW cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;
XX KW cardiovascular disease; peripheral arterial disease; cancer;
XX KW phosphoglycerate kinase; PGK; murine; promoter; OBHrel; ds.
XX OS Mus sp.
XX PN WO200017371-A1.
XX PD 30-MAR-2000.
XX PF 22-SEP-1999; 99WO-GB03181.
XX PR 23-SEP-1998; 98WO-GB02885.
XX PR 28-JAN-1999; 99GB-0001908.
XX PR 16-FEB-1999; 99GB-0003538.
XX PA (OXFO-) OXFORD BIOMEDICA UK LTD.
XX PI Binley KM, Naylor S;
XX DR WPI; 2000-283595/24.
XX PT Novel polynucleotide constructs comprising at least two repeats of a
XX PT hypoxia response element useful for driving expression of nucleic acids
XX PT of interest in a cell under hypoxic conditions
XX PS Example 1; Page 67-68; 155pp; English.
XX CC This invention describes novel polynucleotide comprising at least 2
XX CC repeats of a hypoxia response element (HRE), where the hypoxia-inducible
XX CC factor (HIF) consensus binding sites within each of the 2 repeats are
XX CC separated by a spacer of at least 20 contiguous nucleotides. The products
XX CC of the invention have vasotropic, cardiant, cytostatic and antiarthritic
XX CC activity and can be used for gene therapy. The polynucleotides are useful
XX CC for delivering nucleic acids of interest to mammalian cells. Lentiviral
XX CC vectors are responsive to hypoxic agents and to agents that mimic
XX CC hypoxia. This regulation can be harnessed in vitro to enhance the
XX CC production of the vector and can be used in vivo to regulate gene
XX CC expression in response to a physiological signal. The vectors have
XX CC utility in disease, where ischaemia, including hypoxia, is a feature,
XX CC e.g. cardiovascular disease, peripheral arterial disease, cancer and
XX CC arthritis. The novel regulatory construct is capable of driving very high
XX CC levels of transcription under conditions of hypoxia whilst providing only
XX CC low basal levels of transcription under normal oxygen conditions. The
XX CC polynucleotide construct targets cells within a tumor mass that are under
XX CC conditions of hypoxia without affecting normal surrounding tissue. This
XX CC sequence represents a murine phosphoglycerate kinase (PGK) HRE derived
XX CC promoter OBHrel which is described in the method of the invention.
XX SQ Sequence 243 BP; 45 A; 83 C; 58 G; 57 T; 0 other;

Query Match 93.2%; Score 221; DB 21; Length 243;
Best Local Similarity 97.5%; Pred. No. 5e-58;
Matches 237; Conservative 0; Mismatches 0; Indels 6; Gaps 1;

QY 1 GCTAGAGTCGTGCAGGACGTGACATCTAGTGTGTCGAGG-----CATCTAGTGTCTGG 54
Db 1 GCTAGAGTCGTGCAGGACGTGACATCTAGTGTGTCGAGGACGTGACATCTAGTGTCTGG 60
QY 55 CAGAGCGTGACAGCTAGCGCGGCTCGAGATCTGCATCTGCATCTCAATTAGTCAGCAA 114
Db 61 CAGAGCGTGACAGCTAGCGCGGCTCGAGATCTGCATCTGCATCTCAATTAGTCAGCAA 120
QY 115 CCATAGTCCCGCCCTAACTCGCCCATCCCGCCCTAACTCGCCCATTCGCCCCATT 174

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Db 121 CCATAGTCCCGCCCTAACTCGCCCATCCCGCCCTAACTCGCCCATTCGCCCCATT 180
QY 175 CTCGCCCCATCGCTGACTAATTTTTTTTATTTATGAGAGCGGAGCGCTCGGCCT 234
Db 181 CTCGCCCCATCGCTGACTAATTTTTTTTATTTATGAGAGCGGAGCGCTCGGCCT 240
QY 235 CTG 237
Db 241 CTG 243

RESULT 4
AAZ07789
ID AAZ07789 standard; DNA; 242 BP.
XX AC AAZ07789;
XX DT 23-NOV-1999 (first entry)
XX DE Promoter OBHrel used in EIAV vectors expressing P450.
XX KW Prodrug; localization domain; tumor-selective antibody; cytochrome P450;
XX KW prodrug activating domain; modified hematopoietic stem cell; MHSC; tumor;
XX KW inflammation; atherosclerosis; muscular dystrophy; cerebral malaria; PGK;
XX KW rheumatoid arthritis; hypoxia; ischemia; hypoglycemia; promoter; ss.
XX OS Synthetic.
XX OS Mus sp.
XX OS Rhesus macaque polyoma virus.
XX FH Key Location/Qualifiers
XX FT misc_feature 1..81
XX FT /*tag= a
XX FT /note= "murine PGK"
XX FT promoter 82..214
XX FT /*tag= b
XX FT /note= "SV40 promoter fragment"
XX PN WO9945126-A2.
XX PD 10-SEP-1999.
XX PF 05-MAR-1999; 99WO-GB00672.
XX PR 06-MAR-1998; 98GB-0004841.
XX PR 19-AUG-1998; 98GB-0018103.
XX PR 29-JAN-1999; 99GB-0002081.
XX PA (OXFO-) OXFORD BIOMEDICA UK LTD.
XX PI Stratford IJ, Patterson AV, Kingsman SM, Kan O, Griffiths L;
XX PI Mitrophanous K;
XX DR WPI; 1999-540852/45.
XX PT New prodrug activating agent targeted to selected cells or tissues,
XX PT particularly hypoxic cells, for treating e.g. tumors or inflammation
XX PT Example 14B; Page 97-98; 149pp; English.
XX PS The invention provides a new prodrug activating agent that comprises: (i)
XX CC a localization domain (LD); other than a tumor-selective antibody) and a
XX CC prodrug activating domain (PAD); (ii) at least one nucleic acid encoding
XX CC a cytochrome P450 and under control of at least one constitutive or
XX CC inducible expression control sequence or (iii) a modified hematopoietic
XX CC stem cell (MHSC) containing at least one nucleic acid encoding a PAD and
XX CC under control of elements as in (ii). The prodrug activating agent or
XX CC vectors that express them, are specifically used to treat tumors.
XX CC Inflammation, atherosclerosis and muscular dystrophy, but may also be
XX CC used to treat many other conditions, e.g. cerebral malaria, rheumatoid
XX CC arthritis, or conditions associated with hypoxia, hypoglycemia or
XX CC ischaemia, or to deliver antibiotics, antiviral agents, analgesics,
XX CC anesthetics, anti-inflammatories, antineoplastic agents and diagnostic

```

XX	This invention describes novel polynucleotide comprising at least 2
CC	repeats of a hypoxia response element (HRE), where the hypoxia-inducible
CC	factor (HIF) consensus binding sites within each of the 2 repeats are
CC	separated by a spacer of at least 20 contiguous nucleotides. The products
CC	of the invention have vasotropic, cardiatic, cytostatic and antiarthritic
CC	activity and can be used for gene therapy. The polynucleotides are useful
CC	for delivering nucleic acids of interest to mammalian cells. Lentiviral
CC	vectors are responsive to hypoxic agents and to agents that mimic
CC	hypoxia. This regulation can be harnessed in vitro to enhance the
CC	production of the vector and can be used in vivo to regulate gene
CC	expression in response to a physiological signal. The vectors have
CC	utility in disease, where ischemia, including hypoxia, is a feature,
CC	e.g. cardiovascular disease, peripheral arterial disease, cancer and
CC	arthritis. The novel regulatory construct is capable of driving very high
CC	levels of transcription under conditions of hypoxia whilst providing only
CC	low/basal levels of transcription under normal oxygen conditions. The
CC	polynucleotide construct targets cells within a tumor mass that are under
CC	conditions of hypoxia without affecting normal surrounding tissue. This
CC	sequence represents a murine phosphoglycerate kinase (PGK) HRE derived
CC	promoter element which is described in the method of the invention.
XX	
SQ	Sequence 242 BP; 45 A; 93 C; 47 G; 57 T; 0 other;
	Query Match 73.8%; Score 175; DB 21; Length 242;
	Best Local Similarity 84.8%; Pred. No. 5.9e-44;
	Matches 196; Conservative 0; Mismatches 35; Indels 0; Gaps 0;
QY	7 GTCTGTCAGGCGGTGCACATCTAGTGTGGTGCGAGCATCTAGTCTGTGCAGGACGTGACA 66
DB	12 GHCTCTGCAGCAGCATAGATGTCAGTCTCTGCAGCACACTAGATGTCACGTCTGTGCAGCAG 71
QY	67 GCTAGCCCGGGCTCGAGATCTCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCCG 126
DB	72 TCTAGCCCGGGCTCGAGATCTCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCCG 131
QY	127 CCTTAACCTCGGCCATCCGGCCCCTAACTCGGCCAGTTCGGCCCATTTCTCGGCCCATTC 186
DB	132 CCTTAACCTCGGCCATCCGGCCCCTAACTCGGCCAGTTCGGCCCATTTCTCGGCCCATTC 191
QY	187 GCTGACTAATTTTTTTTATTATGTCAGAGGCGGAGCGCGCTCGGGCTCTG 237
DB	192 GCTGACTAATTTTTTTTATTATGTCAGAGGCGGAGCGCGCTCGGGCTCTG 242
RESULT 6	
AAD27539	
ID	AAD27539 standard; DNA; 5010 BP.
AC	AAD27539;
XX	
DT	18-APR-2002 (first entry)
XX	
DE	pGL3 promoter vector DNA.
XX	
KW	p53 protein; pGL3 luciferase reporter vector; luc+; transcription factor;
KW	cell cycle control; DNA damage repair; pGL3 promoter vector; apoptosis;
KW	firefly; ds.
XX	
OS	Chimeric - Photinus pyralis.
OS	Chimeric - Unidentified.
XX	
FH	Key Location/Qualifiers
FT	misc_feature 1..41
FT	/tag= a
FT	/notes= "Multiple cloning site"
FT	49..244
FT	/tag= b
FT	misc_feature 280..1929
FT	/tag= c
FT	/note= "Luciferase gene (luc+) "
FT	complement (281..303)
FT	/tag= d



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FT polyA_signal /bound_moiety= "GL primer2"
FT 1964..2185 /tag= e
FT /note= "SV40 late poly(A) signal"
FT primer_bind Complement (2253..2272)
FT /tag= f
FT /bound_moiety= "RV primer4"
FT 2510
FT misc_signal /tag= g
FT /note= "ColE1-derived plasmid replication origin"
FT misc_feature Complement (3275..4132)
FT /tag= h
FT /note= "Beta-lactamase gene"
FT misc_signal 4265..4719
FT /tag= i
FT /note= "F1 origin"
FT polyA_signal 4850..5003
FT /tag= j
FT /note= "Upstream poly(A) signal"
FT primer_bind 4952..4971
FT /tag= k
FT /bound_moiety= "RV primer3"
FT WO200196602-A2.
FT XX
FT XX
FT XX
FT XX
FT 18-JUN-2001; 2001WO-GB02718.
FT XX
FT 16-JUN-2000; 2000GB-0014820.
FT PR
FT (MEDI-) MEDICAL RES COUNCIL.
FT PA
FT XX
FT Yang AL, Festing M;
FT PI
FT WPI; 2002-130743/17.
FT DR
FT XX
FT Determining the p53 status of a sample, useful for assaying for
FT mimetics or antagonists of p53, or for the presence of DNA damage,
FT comprises determining whether p53 binds to the pGL3 vector in a sample
FT containing a pGL3 vector -
FT XX
FT Claim 20; Page 42-44; 53pp; English.
FT XX
FT The patent discloses methods for determining the p53 status of a sample
FT which comprise providing a sample containing a pGL3 luciferase reporter
FT vector and determining whether p53 binds to the pGL3 vector. p53 is a
FT transcription factor that regulates many genes including those associated
FT with cell cycle control, apoptosis and DNA damage repair. pGL3 reporter
FT vectors contain a modified firefly luciferase cDNA designated luc+. p53
FT protein binds to pGL3-basic vector and causes luciferase expression. The
FT method is useful for determining the p53 status of a sample. It is also
FT useful for assaying for mimetics or antagonists of p53 and for assaying
FT for presence of activated p53 and/or DNA damage. The invention also
FT relates to a method of modifying pGL3 vector which involves deletion
FT or alteration of a CCGGG motif of the pGL3 vector and/or deleting or
FT altering a sequence within 20 bp sequence 5' or 3' of CCGGG motif. The
FT nucleic acid having a sequence incorporating the CCGGG motif is useful
FT for conferring promoter activity or p53-responsiveness on a nucleic acid
FT encoding a polypeptide of interest or in assays for p53 transcriptional
FT activity. The present DNA sequence is pGL3 promoter vector sequence.
FT XX
FT Sequence 5010 BP; 1287 A; 1213 C; 1198 G; 1312 T; 0 other;
FT SQ
FT
FT Query Match 72.2%; Score 171; DB 24; Length 5010;
FT Best Local Similarity 100.0%; Pred. No. 2.3e-42;
FT Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
FT
FT QY 67 GCTAGCCGGGCTCGAGATCTGCGATCTCAATTAGTCAGCAACCATAGTCCCGC 126
FT |
FT Db 21 GCTAGCCGGGCTCGAGATCTGCGATCTCAATTAGTCAGCAACCATAGTCCCGC 80
FT |
FT QY 127 CCTAACTCCGCCCATCCGCCCTACTCGGCCAGTTCGGCCCAATCTCCGCCCATC 186
FT |

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Db 81 CCTAACTCCGCCCATCCGCCCTACTCGGCCAGTTCGGCCCAATCTCCGCCCATC 140
|
QY 187 GCTGACTAATTTTTTTTATTTATGTCAGAGCGGCGGCTCGGCCTCTG 237
|
Db 141 GCTGACTAATTTTTTTTATTTATGTCAGAGCGGCGGCTCGGCCTCTG 191
|
RESULT 7
AAA07776
ID AAA07776 standard; DNA: 5256 BP.
XX
XX AAA07776;
XX
XX 03-JUL-2000 (first entry)
XX
XX DNA sequence of plasmid pGL2.
XX
XX ced-6; hlced-6; h2ced-6; signal transduction pathway; phagocytosis;
XX cancer; autoimmune disease; neurodegenerative disease; stroke; AIDS;
XX Huntington's disease; myocardial infarction; cytostatic; neuroprotective;
XX cardiant; immunosuppressive; apoptosis modulator; luciferase; ss.
XX
XX Synthetic.
XX
XX WO9964586-A2.
XX
XX 16-DEC-1999.
XX
XX 10-JUN-1999; 99WO-EP04043.
XX
XX 11-JUN-1998; 98GB-0012660.
XX
XX 24-SEP-1998; 98GB-0020816.
XX
XX (DEVG-) DEVGEN NV.
XX
XX Smits E, Van Crieckinge WMR, Bogaert TAOE;
XX
XX WPI; 2000-246285/21.
XX
XX Assays for determining the phagocytosis of apoptotic cells useful for
XX identifying a compound which influences the phagocytic uptake of
XX apoptotic cells and treats cancers and neurodegenerative diseases -
XX
XX Examples; Fig 19; 122pp; English.
XX
XX The invention relates to assays involving two human homologues of
XX Caenorhabditis elegans ced-6 (hlced-6 and h2ced-6) for identifying
XX compounds which function as an inhibitor or an enhancer of a signal
XX transduction pathway. The assays are carried out by measuring
XX phagocytosis of apoptotic cells. The methods are useful for identifying
XX compounds which can act as apoptotic modulators which are useful for
XX treating diseases such as cancer, autoimmune diseases, neurodegenerative
XX diseases such as Huntington's disease, stroke, myocardial infarction and
XX AIDS. The assays are well adapted for medium and high throughput
XX screening using a multi-well plate format. The present sequence
XX represents the DNA sequence of plasmid pGL2 which is suitable for
XX introduction of reporter gene luciferase into Ba/K3 cells.
XX
XX Sequence 5256 BP; 1336 A; 1268 C; 1281 G; 1371 T; 0 other;
XX SQ
XX
XX Query Match 72.2%; Score 171; DB 21; Length 5256;
XX Best Local Similarity 100.0%; Pred. No. 2.3e-42;
XX Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 67 GCTAGCCGGGCTCGAGATCTGCGATCTGCGATCTCAATTAGTCAGCAACCATAGTCCCGC 126
XX |
XX Db 21 GCTAGCCGGGCTCGAGATCTGCGATCTGCGATCTCAATTAGTCAGCAACCATAGTCCCGC 80
XX |
XX QY 127 CCTAACTCCGCCCATCCGCCCTACTCGGCCAGTTCGGCCCAATCTCCGCCCATC 186
XX |
XX Db 81 CCTAACTCCGCCCATCCGCCCTACTCGGCCAGTTCGGCCCAATCTCCGCCCATC 140
XX |

```

QY 187 GCTGACTAAATTTTATTTATTCAGAGCCGAGCGGCGCTCGGCTCTG 237  
 (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||)  
 DB 141 GCTGACTAAATTTTATTTATTCAGAGCCGAGCGGCGCTCGGCTCTG 191

RESULT 8  
 AD27537  
 ID AD27537 standard; DNA; 5256 BP.  
 XX  
 AC AD27537;  
 DT 18-APR-2002 (first entry)  
 XX  
 DE PGL3 control vector DNA.  
 XX  
 KW p53 protein; PGL3 luciferase reporter vector; luc+; transcription factor;  
 KW cell cycle control; DNA damage repair; PGL3 control vector; apoptosis;  
 KW firefly; ds.  
 XX  
 OS Chimeric - Photinus pyralis.  
 OS Chimeric - Unidentified.

XX Key Location/Qualifiers  
 FH 1..41  
 FT misc\_feature /\*tag= a  
 FT /\*note= "Multiple cloning site"  
 FT 49..244 /\*tag= b  
 FT /\*tag= c  
 FT 280..1929 /\*tag= d  
 FT /\*note= "Luciferase gene (luc+)"  
 FT complement (281..303)  
 FT /\*tag= e  
 FT /bound\_moiety= "GL primer2"  
 FT 1964..2185 /\*tag= f  
 FT /\*note= "SV40 late poly(A) signal"  
 FT 2197..2441 /\*tag= g  
 FT complement (2499..2518)  
 FT /\*tag= h  
 FT /bound\_moiety= "RV primer4"  
 FT 2756 /\*tag= i  
 FT /\*note= "ColE1-derived plasmid replication origin"  
 FT complement (3521..4378)  
 FT /\*tag= j  
 FT /\*note= "Beta-lactamase gene"  
 FT 4511..4965 /\*tag= k  
 FT /\*note= "F1 origin"  
 FT 5096..5249 /\*tag= l  
 FT /\*note= "Upstream poly(A) signal"  
 FT 5198..5217 /\*tag= m  
 FT /bound\_moiety= "RV primer3"

XX WO200196602-A2.  
 XX  
 XX 20-DEC-2001.  
 XX  
 XX 18-JUN-2001; 2001WO-GB02718.  
 XX  
 XX 16-JUN-2000; 2000GB-0014820.  
 XX  
 XX (MEDI-) MEDICAL RES COUNCIL.  
 XX  
 XX Yang AL, Festing M;  
 PI  
 XX WPI; 2002-130743/17.  
 DR  
 XX Determining the p53 status of a sample, useful for assaying for

PT mimetics or antagonists of p53, or for the presence of DNA damage,  
 PT comprises determining whether p53 binds to the PGL3 vector in a sample  
 XX containing a PGL3 vector  
 PS Claim 20; Page 36-39; 53pp; English.  
 XX  
 CC The patent discloses methods for determining the p53 status of a sample  
 CC which comprise providing a sample containing a PGL3 luciferase reporter  
 CC vector and determining whether p53 binds to the PGL3 vector. p53 is a  
 CC transcription factor that regulates many genes including those associated  
 CC with cell cycle control, apoptosis and DNA damage repair. PGL3 reporter  
 CC vectors contain a modified firefly luciferase cDNA designated luc+. p53  
 CC protein binds to PGL3-basic vector and causes luciferase expression. The  
 CC method is useful for determining the p53 status of a sample. It is also  
 CC useful for assaying for mimetics or antagonists of p53 and for assaying  
 CC for presence of activated p53 and/or DNA damage. The invention also  
 CC relates to a method of modifying PGL3 vector which involves deleting or  
 CC or alteration of a CCGGG motif of the PGL3 vector and/or deleting or  
 CC altering a sequence within 20 bp sequence 5' or 3' of CCGGG motif. The  
 CC nucleic acid having a sequence incorporating the CCGGG motif is useful  
 CC for conferring promoter activity or p53-responsiveness on a nucleic acid  
 CC encoding a polypeptide of interest or in assays for p53 transcriptional  
 CC activity. The present DNA sequence is PGL3 control vector sequence.

XX Sequence 5256 BP; 1336 A; 1268 C; 1281 G; 1371 T; 0 other;  
 SQ  
 Query Match 72.2%; Score 171; DB 24; Length 5256;  
 Best Local Similarity 100.0%; Pred. No. 2.3e-42;  
 Matches 171; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 67 GCTAGCCGGGCTCGAGATCTGCGATCTGCAATTCATTCAGCAACCATAGTCCCGC 126  
 |||||  
 DB 21 GCTAGCCGGGCTCGAGATCTGCGATCTGCAATTCATTCAGCAACCATAGTCCCGC 80  
 QY 127 CCTAACTCCGCCATCCGCCCTAACTCCGCCATTCGCCGCCATTCGCCGCCATC 186  
 |||||  
 DB 81 CCTAACTCCGCCATCCGCCCTAACTCCGCCATTCGCCGCCATTCGCCGCCATC 140  
 QY 187 GCTGACTAAATTTTATTTATTCAGAGCCGAGCGGCGCTCGGCTCTG 237  
 |||||  
 DB 141 GCTGACTAAATTTTATTTATTCAGAGCCGAGCGGCGCTCGGCTCTG 191

RESULT 9  
 AAAL2019  
 ID AAAL2019 standard; DNA; 267 BP.  
 XX  
 AC AAAL2019;  
 XX  
 DT 14-AUG-2000 (first entry)  
 XX  
 DE Human erythropoietin enhancer Epo X4 DNA.  
 XX  
 KW HFE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;  
 KW cardiac; cytosolic; antiarthritic; gene therapy; ischaemia; arthritis;  
 KW cardiovascular disease; peripheral arterial disease; cancer; human;  
 KW enhancer; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200017371-A1.  
 XX  
 XX 30-MAR-2000.  
 XX  
 XX 22-SEP-1999; 99WO-GB03181.  
 XX  
 XX 23-SEP-1998; 98WO-GB02885.  
 XX 28-JAN-1999; 99GB-0001906.  
 XX 16-FEB-1999; 99GB-0003538.  
 XX  
 XX (OXFO-) OXFORD BIOMEDICA UK LTD.  
 XX  
 XX Binley KM, Naylor S;  
 PI

```

XX WPI; 2000-283595/24.
XX
XX Novel polynucleotide constructs comprising at least two repeats of a
XX hypoxia response element useful for driving expression of nucleic acids
XX of interest in a cell under hypoxic conditions
XX
XX Example 1; Page 69; 155pp; English.
XX
XX This invention describes novel polynucleotide comprising at least 2
XX repeats of a hypoxia response element (HRE), where the hypoxia-inducible
XX factor (HIF) consensus binding sites within each of the 2 repeats are
XX separated by a spacer of at least 20 contiguous nucleotides. The products
XX of the invention have vasotropic, cardiant, cytostatic and antiarthritic
XX activity and can be used for gene therapy. The polynucleotides are useful
XX for delivering nucleic acids of interest to mammalian cells. Lentiviral
XX vectors are responsive to hypoxic agents and to agents that mimic
XX hypoxia. This regulation can be harnessed in vitro to enhance the
XX production of the vector and can be used in vivo to regulate gene
XX expression in response to a physiological signal. The vectors have
XX utility in disease, where ischaemia, including hypoxia, is a feature,
XX e.g. cardiovascular disease, peripheral arterial disease, cancer and
XX arthritis. The novel regulatory construct is capable of driving very high
XX levels of transcription under conditions of hypoxia whilst providing only
XX low basal levels of transcription under normal oxygen conditions. The
XX polynucleotide construct targets cells within a tumor mass that are under
XX conditions of hypoxia without affecting normal surrounding tissue. This
XX sequence represents a murine phosphoglycerate kinase (PGK) HRE derived
XX promoter element which is described in the method of the invention.
XX
XX Sequence 267 BP; 44 A; 106 C; 52 G; 65 T; 0 other;
XX
XX Query Match 64.2%; Score 152.2; DB 21; Length 267;
XX Best Local Similarity 79.0%; Pred. No. 5.7e-37;
XX Matches 181; Conservative 0; Mismatches 48; Indels 0; Gaps 0;
XX
XX QY 9 CGTCGAGCAGCTGACATCTAGTGTCTGTCAGGCAATCTCTCTGTGAGAGGTGAGC 68
XX Db ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
XX
XX QY 39 CGTGTCTCTACACAGCCTGGATCTGCCCTTACGTGTCTCTACACAGGCTGGCCCTAC 98
XX Db ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
XX
XX QY 69 TAGCCGGGCTCGAGATCTGCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCGCC 128
XX Db || || || || || || || || || || || || || || || || || || ||
XX
XX QY 99 GTGCTGTCTCACAGAGCTGGGATCTGCATCTCAATTAGTCAGCAACCATAGTCCGCC 158
XX Db || || || || || || || || || || || || || || || || || || ||
XX
XX QY 129 CTAAGTCGCGCCATCCGCCCTTAACCTCGGCCAGTTCGCCCATCTCTCGGCCCATCGC 188
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 159 CTAAGTCGCGCCATCCGCCCTTAACCTCGGCCAGTTCGCCCATCTCTCGGCCCATCGC 218
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 189 TGACTAATTTTTTTTATTTATTCAGAGGCGGAGCGCGCTCGGCCCTCTG 237
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 219 TGACTAATTTTTTTTATTTATTCAGAGGCGGAGCGCGCTCGGCCCTCTG 267
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX RESULT 10
XX AAA12021
XX ID AAA12021 standard; DNA; 204 BP.
XX AC AAA12021;
XX XX
XX 14-AUG-2000 (first entry)
XX
XX Murine lactate dehydrogenase A regulatory element DNA.
XX
XX HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;
XX cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;
XX cardiovascular disease; peripheral arterial disease; cancer;
XX lactate dehydrogenase A; murine; ds.
XX
XX Mus sp.
XX
XX WO200017371-A1.
XX
XX 30-MAR-2000.
XX

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XX 22-SEP-1999; 99WO-GB03181.
XX
XX 23-SEP-1998; 98WO-GB02885.
XX PR 28-JAN-1999; 99GB-0001906.
XX PR 16-FEB-1999; 99GB-0003538.
XX
XX (OXFO-) OXFORD BIOMEDICA UK LTD.
XX
XX Binley KM, Naylor S;
XX WPI; 2000-283595/24.
XX
XX Novel polynucleotide constructs comprising at least two repeats of a
XX hypoxia response element useful for driving expression of nucleic acids
XX of interest in a cell under hypoxic conditions
XX
XX Example 1; Page 70; 155pp; English.
XX
XX This invention describes novel polynucleotide comprising at least 2
XX repeats of a hypoxia response element (HRE), where the hypoxia-inducible
XX factor (HIF) consensus binding sites within each of the 2 repeats are
XX separated by a spacer of at least 20 contiguous nucleotides. The products
XX of the invention have vasotropic, cardiant, cytostatic and antiarthritic
XX activity and can be used for gene therapy. The polynucleotides are useful
XX for delivering nucleic acids of interest to mammalian cells. Lentiviral
XX vectors are responsive to hypoxic agents and to agents that mimic
XX hypoxia. This regulation can be harnessed in vitro to enhance the
XX production of the vector and can be used in vivo to regulate gene
XX expression in response to a physiological signal. The vectors have
XX utility in disease, where ischaemia, including hypoxia, is a feature,
XX e.g. cardiovascular disease, peripheral arterial disease, cancer and
XX arthritis. The novel regulatory construct is capable of driving very high
XX levels of transcription under conditions of hypoxia whilst providing only
XX low basal levels of transcription under normal oxygen conditions. The
XX polynucleotide construct targets cells within a tumor mass that are under
XX conditions of hypoxia without affecting normal surrounding tissue. This
XX sequence represents a murine lactate dehydrogenase A HRE regulatory
XX element which is described in the method of the invention.
XX
XX Sequence 204 BP; 33 A; 81 C; 41 G; 49 T; 0 other;
XX
XX Query Match 62.6%; Score 148.4; DB 21; Length 204;
XX Best Local Similarity 99.3%; Pred. No. 7.7e-36;
XX Matches 149; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
XX QY 88 GCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCGCCCATCTCGGCCCATCTCCGC 147
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 55 GGGATCTGCATCTCAATTAGTCAGCAACCATAGTCCGCCCATCTCGGCCCATCTCCGC 114
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 148 CCCTAACTCCGCCAGTTCGCCCATCTCGGCCCATCTCGGCCCATCTCGGCCCATCTTT 207
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 115 CCCTAACTCCGCCAGTTCGCCCATCTCGGCCCATCTCGGCCCATCTCGGCCCATCTTT 174
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 208 ATGCAGAGCGGAGCGCGCTCGGCCCTCTG 237
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX QY 175 ATGCAGAGCGGAGCGCGCTCGGCCCTCTG 204
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
XX RESULT 11
XX AAA12017
XX ID AAA12017 standard; DNA; 223 BP.
XX AC AAA12017;
XX XX
XX 14-AUG-2000 (first entry)
XX
XX Human enolase A promoter element DNA.
XX
XX HRE; hypoxia response element; hypoxia-inducible factor; HIF; vasotropic;
XX cardiant; cytostatic; antiarthritic; gene therapy; ischaemia; arthritis;
XX cardiovascular disease; peripheral arterial disease; cancer; human;
XX promoter; ds.
XX

```

X	Homo sapiens.
S	WC200017371-A1.
X	N
N	D
X	30-MAR-2000.
X	22-SEP-1999; 99WO-GB03181.
X	F
X	23-SEP-1998; 98WO-GB02885.
R	R
R	28-JAN-1999; 99GB-0001906.
R	R
R	16-FEB-1999; 99GB-0003538.
A	(OXFO-) OXFORD BIOMEDICA UK LTD.
X	Binley KM, Naylor S;
X	WPI; 2000-283595/24.
I	Novel polynucleotide constructs comprising at least two repeats of a
X	hypoxia response element useful for driving expression of nucleic acids
T	of interest in a cell under hypoxic conditions -
T	Example 1; Page 69; 155pp; English.
S	This invention describes novel polynucleotide comprising at least 2
X	repeats of a hypoxia response element (HRE), where the hypoxia-inducible
C	factor (HIF) consensus binding sites within each of the 2 repeats are
C	separated by a spacer of at least 20 contiguous nucleotides. The products
C	of the invention have vasotropic, cardiant, cytostatic and antiarthritic
C	activity and can be used for gene therapy. The polynucleotides are useful
C	for delivering nucleic acids of interest to mammalian cells. Lentiviral
C	vectors are responsive to hypoxic agents and to agents that mimic
C	hypoxia. This regulation can be harnessed in vitro to enhance the
C	production of the vector and can be used in vivo to regulate gene
C	expression in response to a physiological signal. The vectors have
C	utility in disease, where ischaemia, including hypoxia, is a feature,
C	e.g. cardiovascular disease, peripheral arterial disease, cancer and
C	arthritis. The novel regulatory construct is capable of driving very high
C	levels of transcription under conditions of hypoxia whilst providing only
C	low basal levels of transcription under normal oxygen conditions. The
C	polynucleotide construct targets cells within a tumor mass that are under
C	conditions of hypoxia without affecting normal surrounding tissue. This
C	sequence represents a murine phosphoglycerate kinase (pgk) HRE derived
C	promoter element which is described in the method of the invention.
X	Sequence 223 BP; 38 A; 81 C; 58 G; 46 T; 0 other;
X	Query Match 62.4%; Score 148; DB 21; Length 223;
SQ	Best Local Similarity 100.0%; Pred. No. 1e-35;
	Matches 148; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	90 GATCTGCATCAATTAGTCAGACACCATTGTCGCCGCCCTAACTCCGCCCATCCGGCCC 149
DB	76 GATCTGCATCTCAATTAGTCAGACACCATTGTCGCCGCCCTAACTCCGCCCATCCGGCCC 135
QY	150 CTAACTCCGCCCGAGTTCGCCCATTTCTCGGCCCATTCGCTGACTAATTTTTTTTAT 209
DB	136 CTAACTCCGCCCGAGTTCGCCCATTTCTCGGCCCATTCGCTGACTAATTTTTTTTAT 195
QY	210 GCAGAGCGCGAGCGCGCTCGGCTCTG 237
DB	196 GCAGAGCGCGAGCGCGCTCGGCTCTG 223
RESULT 12	
AA42469	
ID	AA42469 standard; cDNA; 356 BP.
XX	AC
XX	AA42469;
XX	05-AUG-1997 (first entry)

DE	Interleukin-6 inhibitor BamHI-HindIII fragment of pM8SV.
XX	inhibitor; interleukin-6; IL-6; acute phase response element; APRE;
KW	transcription factor binding site; therapy; autoimmune; Castleman's;
KW	inflammatory disease; rheumatoid arthritis; glomerulonephritis;
KW	psoriasis; ss.
XX	Synthetic.
OS	
XX	Location/Qualifiers
FH	Key 11..155
FT	misc_feature /*tag= a
FT	/note= "M8 sequence"
FT	11..19
FT	misc_signal /*tag= b
FT	/note= "acute phase response element; claim 2"
FT	160..356
FT	misc_feature /*tag= C
FT	/note= "SV40 promoter sequence"
XX	
PN	WO9635782-A1.
PX	
PD	14-NOV-1996.
XX	
PF	11-MAY-1995; 95WO-EP01778.
XX	
PR	11-MAY-1995; 95WO-EP01778.
XX	
PA	(ISTF ) ARS APPLIED RES SYST HOLDING NV.
XX	
PI	Pezzotti A, Serlupi-Crescenzi O;
XX	WPI; 1996-518670/51.
DR	
XX	Nucleotide sequence which inhibits interleukin-6 activity - used in
PT	pharmaceutical compositions for treatment of auto-immune and
PT	inflammatory diseases
XX	Claim 10; Fig 2; 35pp; English.
PS	
XX	The present sequence comprises the BamHI-HindIII fragment of pM8SVL
CC	which contains a nucleotide sequence which is able to inhibit
CC	interleukin-6 (IL-6) activity. The nucleotide sequence contains an acute
CC	phase response element (APRE) and at least one nucleotide sequence
CC	constituting a transcription factor binding site other than the APRE
CC	element. The nucleotide sequences can be used in therapy to inhibit the
CC	action of IL-6 in conditions where IL-6 plays a pathological role, e.g.
CC	autoimmune and inflammatory diseases such as rheumatoid arthritis,
CC	psoriasis, Castleman's disease and glomerulonephritis. Other methods of
CC	inhibiting IL-6 activity, such as use of antibodies against IL-6, gp130
CC	or gp80, or the use of mutants for IL-6 or the IL-6 receptor, may be
CC	associated with undesired clinical effects. The new approach involves
CC	blocking the intracellular proteins mediating the IL-6 signal.
XX	
SQ	Sequence 356 BP; 66 A; 103 C; 76 G; 111 T; 0 other;
	Query Match 62.2%; Score 147.4; DB 17; Length 356;
	Best Local Similarity 96.2%; Pred. No. 1.8e-35;
	Matches 151; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
QY	81 GAGATCTGGCATCTGCATCTCAATTAGTCAGACACCATTGTCGCCGCCCTAACTCCGCCCC 140
DB	141 GGAATTCTGTATCTGCATCTCAATTAGTCAGACACCATTGTCGCCGCCCTAACTCCGCCCC 200
QY	141 ATCCCGCCCCTAACCTCCGCCCAGTTCGCCCATTTCTCGGCCCATTCGCTGACTAATTTTT 200
DB	201 ATCCCGCCCCTAACCTCCGCCCAGTTCGCCCATTTCTCGGCCCATTCGCTGACTAATTTTT 260
QY	201 TTATTTATGACAGAGCGCGAGCGCGCTCGGCCTCTG 237
DB	261 TTATTTATGACAGAGCGCGAGCGCGCTCGGCCTCTG 297

RESULT 13  
 AAX08776  
 ID AAX08776 standard; DNA; 5793 BP.  
 AC AAX08776;  
 DT 27-SEP-1999 (first entry)  
 XX pGL2(apo AI-ARE)SV40/luc plasmid construct comprising ARE/luciferase.  
 DE Antioxidant responsive element; ARE; low density lipoprotein; LDL;  
 KW high density lipoprotein; HDL; apolipoprotein; apo AI;  
 KW atherosclerosis; heart disease; transcription; ss.  
 XX Synthetic.  
 XX CA2238662-A.  
 PN 23-NOV-1998.  
 XX 22-MAY-1998; 98CA-2238662.  
 PF 23-MAY-1997; 97US-0862431.  
 PR (TOOH ) UNIV QUEENS KINGSTON.  
 PA Tam S;  
 PI WPI; 1999-229918/20.  
 DR New Antioxidant Response Element (ARE), useful for identifying drugs  
 PT and transcription factors for increasing transcription of mRNA,  
 PT useful for treatment of atherosclerosis  
 XX Examples; Page 63-66; 115pp; English.  
 XX DNA constructs comprising antioxidant responsive elements (AREs)  
 CC are useful for screening for compounds and transcription factors  
 CC that bind to the ARE and increase transcription levels of a mRNA  
 CC regulated by an ARE. AREs may also be useful as a reagent for  
 CC purification of a compound (preferably a transcription factor)  
 CC with which it interacts. High Density Lipoprotein (HDL) has  
 CC antioxidant activity and protects against oxidized low-density  
 CC lipoprotein (LDL) which has a role in the etiology of  
 CC atherosclerosis. Apolipoprotein (apo) AI is a major component of  
 CC HDL, and is believed to promote the process of reverse cholesterol  
 CC transport. The transcription of apo AI is effected by cis- and  
 CC trans-acting factors (i.e an ARE) UV cross-linking studies using  
 CC an apoAI-ARE probe isolated two polypeptides of 100 and 115 kDa.  
 CC These compounds are useful for treatment of a human or animal  
 CC with atherosclerosis. ARE's can also be used in DNA constructs  
 CC effect the transcription of those heterologous sequences. A  
 CC Genelight vector (pGL2-B) comprising a 491 bp fragment of the apo AI  
 CC from -491 to +1 inserted into the XhoI site of the vector and  
 CC upstream of the luciferase gene was designated pGL2(apoAI-491)luc.  
 CC Plasmid pGL2(apoAI-250)luc was constructed by releasing a DNA  
 CC fragment (-491 to -251 of the apo AI promoter) from  
 CC pGL2(apoAI-491)luc. A new plasmid pGL2(apoAI-250 mutant ARE) was  
 CC prepared by PCR using four primers (AAX08785-88). Two primers  
 CC (AAX08785, AAX08786) were complementary to the non-coding strand of DNA,  
 CC the remaining two primers (AAX08787, AAX08788) were complementary to  
 CC the coding strand of DNA. The plasmid pGL2(apoAI-250)luc was used  
 CC as a template and the DNA fragment was purified and then cloned into  
 CC the SmaI and HindIII sites of pGL2-B to generate  
 CC pGL2(apoAI-250 mutant ARE). A series of pGL2-Promoter vectors  
 CC comprising apoAI-ARE, GST-ARE and mutated ARE were also constructed.  
 CC See AAX08773-X08779.  
 XX Sequence 5793 BP; 1562 A; 1330 C; 1310 G; 1591 T; 0 other;  
 SQ Query Match 62.08; Score 147; DB 20; Length 5793;  
 Best Local Similarity 86.68; Pred. No. 5.3e-35;

Matches 162; Conservative 0; Mismatches 25; Indels 0; Gaps 0;  
 QY 51 CGTCGAGGACGTCGACAGCTAGCCCGGGCTCGAGATCTGCGATCTCAATTAGTCA 110  
 DB 3 CGGAGGTACACAGCCCGGAGGACAGAGCTGCTAGCTCGAGATCTCAATTAGTCA 62  
 QY 111 GCAACATAGTCCCGCCCTAACTCCGCCATCCGCCCTAACTCCGCCAGTTCGGCC 170  
 DB 63 GCAACATAGTCCCGCCCTAACTCCGCCATCCGCCCTAACTCCGCCAGTTCGGCC 122  
 QY 171 CATTCTCGCCCGCTGCTGACTAATTTTATTCAGAGCGCGAGCGGCTCG 230  
 DB 123 CATTCTCGCCCGCTGCTGACTAATTTTATTCAGAGCGCGAGCGGCTCG 182  
 QY 231 GCCTCTG 237  
 DB 183 GCCTCTG 189  
 RESULT 14  
 AAX08779  
 ID AAX08779 standard; DNA; 5789 BP.  
 XX AAX08779;  
 XX 27-SEP-1999 (first entry)  
 DT pGL2-Promoter genelight vector comprising ARE and apo-AI promoter.  
 DE Antioxidant responsive element; ARE; low density lipoprotein; LDL;  
 KW high density lipoprotein; HDL; apolipoprotein; apo AI;  
 KW atherosclerosis; heart disease; transcription; ss.  
 XX Synthetic.  
 OS CA2238662-A.  
 PN 23-NOV-1998.  
 PD 22-MAY-1998; 98CA-2238662.  
 PF 23-MAY-1997; 97US-0862431.  
 PR (TOOH ) UNIV QUEENS KINGSTON.  
 PA Tam S;  
 PI WPI; 1999-229918/20.  
 DR New Antioxidant Response Element (ARE), useful for identifying drugs  
 PT and transcription factors for increasing transcription of mRNA,  
 PT useful for treatment of atherosclerosis  
 XX Examples; Page 74-77; 115pp; English.  
 XX DNA constructs comprising antioxidant responsive elements (AREs)  
 CC are useful for screening for compounds and transcription factors  
 CC that bind to the ARE and increase transcription levels of a mRNA  
 CC regulated by an ARE. AREs may also be useful as a reagent for  
 CC purification of a compound (preferably a transcription factor)  
 CC with which it interacts. High Density Lipoprotein (HDL) has  
 CC antioxidant activity and protects against oxidized low-density  
 CC lipoprotein (LDL) which has a role in the etiology of  
 CC atherosclerosis. Apolipoprotein (apo) AI is a major component of  
 CC HDL, and is believed to promote the process of reverse cholesterol  
 CC transport. The transcription of apo AI is effected by cis- and  
 CC trans-acting factors (i.e an ARE) UV cross-linking studies using  
 CC an apoAI-ARE probe isolated two polypeptides of 100 and 115 kDa.  
 CC These compounds are useful for treatment of a human or animal  
 CC with atherosclerosis. ARE's can also be used in DNA constructs  
 CC effect the transcription of those heterologous sequences. A  
 CC Genelight vector (pGL2-B) comprising a 491 bp fragment of the apo AI  
 CC from -491 to +1 inserted into the XhoI site of the vector and  
 CC upstream of the luciferase gene was designated pGL2(apoAI-491)luc.  
 CC Plasmid pGL2(apoAI-250)luc was constructed by releasing a DNA  
 CC fragment (-491 to -251 of the apo AI promoter) from  
 CC pGL2(apoAI-491)luc. A new plasmid pGL2(apoAI-250 mutant ARE) was  
 CC prepared by PCR using four primers (AAX08785-88). Two primers  
 CC (AAX08785, AAX08786) were complementary to the non-coding strand of DNA,  
 CC the remaining two primers (AAX08787, AAX08788) were complementary to  
 CC the coding strand of DNA. The plasmid pGL2(apoAI-250)luc was used  
 CC as a template and the DNA fragment was purified and then cloned into  
 CC the SmaI and HindIII sites of pGL2-B to generate  
 CC pGL2(apoAI-250 mutant ARE). A series of pGL2-Promoter vectors  
 CC comprising apoAI-ARE, GST-ARE and mutated ARE were also constructed.  
 CC See AAX08773-X08779.  
 XX Sequence 5793 BP; 1562 A; 1330 C; 1310 G; 1591 T; 0 other;  
 SQ Query Match 62.08; Score 147; DB 20; Length 5793;  
 Best Local Similarity 86.68; Pred. No. 5.3e-35;

regulated by an ARE. AREs may also be useful as a reagent for purification of a compound (preferably a transcription factor) with which it interacts. High Density Lipoprotein (HDL) has antioxidant activity and protects against oxidized low-density lipoprotein (LDL) which has a role in the etiology of atherosclerosis. Apolipoprotein (apo) AI is a major component of HDL, and is believed to promote the process of reverse cholesterol transport. The transcription of apo AI is effected by cis- and trans-acting factors (i.e. an ARE) UV cross-linking studies using an apoAI-ARE probe isolated two polypeptides of 100 and 115 kDa. These compounds are useful for treatment of a human or animal with atherosclerosis. ARE's can also be used in DNA constructs when operably linked to heterologous protein coding sequences. A gene/light vector (pGL2-B) comprising a 491 bp fragment of the apo AI from -491 to +1 inserted into the XhoI site of the vector and upstream of the luciferase gene was designated pGL2(apoAI-491)luc. Plasmid pGL2(apoAI-250)luc was constructed by releasing a DNA fragment (-491 to -251 of the apo AI promoter) from pGL2(apoAI-491)luc. A new plasmid pGL2(apoAI-250 mutant ARE) was prepared by PCR using four primers (AAx08785-88). Two primers (AAx08785, AAx08786) were complementary to the non-coding strand of DNA, the remaining two primers (AAx08787, AAx08788) were complementary to the coding strand of DNA. The plasmid pGL2(apoAI-250)luc was used as a template and the DNA fragment was purified and then cloned into the SmaI and HindIII sites of pGL2-B to generate pGL2(apoAI-250 mutant ARE). A series of pGL2-Promoter vectors comprising apoAI-ARE, GST-ARE and mutated ARE were also constructed. See AAx08773-X08779.

Sequence 5789 BP; 1559 A; 1328 C; 1308 G; 1594 T; 0 other;

Query Match 61.9%; Score 146.8; DB 20; Length 5789;  
Best Local Similarity 98.7%; Pred. No. 6.1e-35;  
Matches 148; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 88 GCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCCGC 147  
DB 36 GAGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCCGC 95  
QY 148 CCTAACTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCT 207  
DB 96 CCTAACTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCT 155  
QY 208 ATGCAGAGCGCGAGCGCGCTCGGCTCTG 237  
DB 156 ATGCAGAGCGCGAGCGCGCTCGGCTCTG 185

RESULT 15  
AAx08777  
ID AAx08777 standard; DNA; 5793 BP.  
AC AAx08777;  
DT 27-SEP-1999 (first entry)  
DE pGL2(apo AI-mARE)SV40/luc construct comprising ARE/luciferase.  
KW Antioxidant responsive element; ARE; low density lipoprotein; LDL;  
KW high density lipoprotein; HDL; apolipoprotein; apo AI;  
KW atherosclerosis; heart disease; transcription; ss.  
XX Synthetic.  
XX CA2238662-A.  
XX 23-NOV-1998.  
XX 22-MAY-1998; 98CA-2238662.  
XX 23-MAY-1997; 97US-0862431.  
XX (TOOH ) UNIV QUEENS KINGSTON.  
XX Tam S;  
XX WPI; 1999-229918/20.  
XX New Antioxidant Response Element (ARE), useful for identifying drugs  
PT and transcription factors for increasing transcription of mRNA,  
PT useful for treatment of atherosclerosis  
XX Examples; Page 66-70; 115pp; English.  
PS DNA constructs comprising antioxidant responsive elements (AREs)  
CC are useful for screening for compounds and transcription factors  
CC that bind to the ARE and increase transcription levels of a mRNA

regulated by an ARE. AREs may also be useful as a reagent for purification of a compound (preferably a transcription factor) with which it interacts. High Density Lipoprotein (HDL) has antioxidant activity and protects against oxidized low-density lipoprotein (LDL) which has a role in the etiology of atherosclerosis. Apolipoprotein (apo) AI is a major component of HDL, and is believed to promote the process of reverse cholesterol transport. The transcription of apo AI is effected by cis- and trans-acting factors (i.e. an ARE) UV cross-linking studies using an apoAI-ARE probe isolated two polypeptides of 100 and 115 kDa. These compounds are useful for treatment of a human or animal with atherosclerosis. ARE's can also be used in DNA constructs when operably linked to heterologous protein coding sequences. A gene/light vector (pGL2-B) comprising a 491 bp fragment of the apo AI from -491 to +1 inserted into the XhoI site of the vector and upstream of the luciferase gene was designated pGL2(apoAI-491)luc. Plasmid pGL2(apoAI-250)luc was constructed by releasing a DNA fragment (-491 to -251 of the apo AI promoter) from pGL2(apoAI-491)luc. A new plasmid pGL2(apoAI-250 mutant ARE) was prepared by PCR using four primers (AAx08785-88). Two primers (AAx08785, AAx08786) were complementary to the non-coding strand of DNA, the remaining two primers (AAx08787, AAx08788) were complementary to the coding strand of DNA. The plasmid pGL2(apoAI-250)luc was used as a template and the DNA fragment was purified and then cloned into the SmaI and HindIII sites of pGL2-B to generate pGL2(apoAI-250 mutant ARE). A series of pGL2-Promoter vectors comprising apoAI-ARE, GST-ARE and mutated ARE were also constructed. See AAx08773-X08779.

Sequence 5793 BP; 1561 A; 1328 C; 1308 G; 1596 T; 0 other;

Query Match 61.9%; Score 146.8; DB 20; Length 5793;  
Best Local Similarity 98.7%; Pred. No. 6.1e-35;  
Matches 148; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 88 GCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCCGC 147  
DB 40 GAGATCTGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCCGC 99  
QY 148 CCTAACTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCT 207  
DB 100 CCTAACTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCT 159  
QY 208 ATGCAGAGCGCGAGCGCGCTCGGCTCTG 237  
DB 160 ATGCAGAGCGCGAGCGCGCTCGGCTCTG 189

Search completed: January 3, 2003, 23:20:43  
Job time : 398.246 secs





Mon Jan 6 15:20:25 2003

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; Patent No. 6120994
; GENERAL INFORMATION:
; APPLICANT: TAM, SHUI-PANG
; TITLE OF INVENTION: ANTIOXIDANT RESPONSIVE ELEMENT
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 NEW YORK AVENUE, SUITE 600
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: US
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/862,431
; FILING DATE: 23-MAY-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Kim, Judith U.
; REGISTRATION NUMBER: 40,679
; REFERENCE/DOCKET NUMBER: 1669.0020000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 371-2600
; TELEFAX: (202) 371-2540
; INFORMATION FOR SEQ ID NO: 32:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 5789 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; US-08-862-431-32

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; Patent No. 6120994
; GENERAL INFORMATION:
; APPLICANT: TAM, SHUI-PANG
; TITLE OF INVENTION: ANTIOXIDANT RESPONSIVE ELEMENT
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 NEW YORK AVENUE, SUITE 600
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: US
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/862,431
; FILING DATE: 23-MAY-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Kim, Judith U.
; REGISTRATION NUMBER: 40,679
; REFERENCE/DOCKET NUMBER: 1669.0020000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 371-2600
; TELEFAX: (202) 371-2540
; INFORMATION FOR SEQ ID NO: 32:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 5789 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; US-08-862-431-32

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Query Match 61.9%; Score 146.8; DB 3; Length 5789;
Best Local Similarity 98.7%; Pred. No. 1.8e-35;
Matches 148; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 88 GCGATCTGCATCTCAATTAGTCAGCAACCATAGTCCGCCCTAACTCCGCCCATCCGC 147
DB 36 GAGATCTGCATCTCAATTAGTCAGCAACCATAGTCCGCCCTAACTCCGCCCATCCGC 95
QY 148 CCTAACTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCAT 207
DB 96 CCTAACTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCATCTCCGCCCAT 155
QY 208 ATGCAGAGCGCGAGCGCGCTCGGCGCTCTG 237
DB 156 ATGCAGAGCGCGAGCGCGCTCGGCGCTCTG 185

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RESULT 4
US-08-862-431-31
; Sequence 31, Application US/08862431
; Patent No. 6120994
; GENERAL INFORMATION:
; APPLICANT: TAM, SHUI-PANG
; TITLE OF INVENTION: ANTIOXIDANT RESPONSIVE ELEMENT
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 NEW YORK AVENUE, SUITE 600
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: US
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30

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QY 141 ATCCGCCCTTAATCCGCCAGTCCGCCATCTCCGCCCATCTCCGCCCATCTTAATTTT 200
DB 201 ATCCGCCCTTAATCCGCCAGTCCGCCATCTCCGCCCATCTCCGCCCATCTTAATTTT 260
QY 201 TTTATTTATGAGAGCGCGAGCGCGCTCGGCGCTCTG 237
DB 261 TTTATTTATGAGAGCGCGAGCGCGCTCGGCGCTCTG 297

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; Patent No. 6120994
; GENERAL INFORMATION:
; APPLICANT: TAM, SHUI-PANG
; TITLE OF INVENTION: ANTIOXIDANT RESPONSIVE ELEMENT
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 NEW YORK AVENUE, SUITE 600
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: US
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/862,431
; FILING DATE: 23-MAY-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Kim, Judith U.
; REGISTRATION NUMBER: 40,679
; REFERENCE/DOCKET NUMBER: 1669.0020000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 371-2600
; TELEFAX: (202) 371-2540
; INFORMATION FOR SEQ ID NO: 29:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 5793 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; US-08-862-431-29

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; Patent No. 6120994
; GENERAL INFORMATION:
; APPLICANT: TAM, SHUI-PANG
; TITLE OF INVENTION: ANTIOXIDANT RESPONSIVE ELEMENT
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 NEW YORK AVENUE, SUITE 600
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: US
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/862,431
; FILING DATE: 23-MAY-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Kim, Judith U.
; REGISTRATION NUMBER: 40,679
; REFERENCE/DOCKET NUMBER: 1669.0020000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 371-2600
; TELEFAX: (202) 371-2540
; INFORMATION FOR SEQ ID NO: 29:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 5793 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; US-08-862-431-29

```

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Query Match 62.0%; Score 147; DB 3; Length 5793;
Best Local Similarity 86.6%; Pred. No. 1.5e-35;
Matches 162; Conservative 0; Mismatches 25; Indels 0; Gaps 0;

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QY 51 GGTGAGAGCTGACAGTCTAGCGCGGCTCGAGATCTGCGATCTGCAATTAATGCA 110
DB 3 CGGAGGTACAGCCCGGAGGACAGAGCTGCTAGCTGAGATCTGCAATTAATGCA 62
QY 111 GCAACCATAGTCCGCCCGCTTAATCTCCGCCCATCTCCGCCCATCTCCGCCCAT 170
DB 63 GCAACCATAGTCCGCCCGCTTAATCTCCGCCCATCTCCGCCCATCTCCGCCCAT 122
QY 171 CATTCCTCCGCCCATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 230
DB 123 CATTCCTCCGCCCATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 182
QY 231 GCTCTG 237
DB 183 GCTCTG 189

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RESULT 3
US-08-862-431-32
; Sequence 32, Application US/08862431

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RESULT 7  
US-09-301-593-42  
; Sequence 42, Application US/09301593A  
; Patent No. 6455677  
; GENERAL INFORMATION:  
; APPLICANT: Park, John E.



;; TITLE OF INVENTION: HUMAN NEUTRALIZING MONOCLONAL ANTIBODIES  
;; TITLE OF INVENTION: TO HUMAN IMMUNODEFICIENCY VIRUS  
;; NUMBER OF SEQUENCES: 170  
;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: The Scripps Research Institute, Office of  
;; ADDRESSEE: Patent Counsel  
;; STREET: 10666 No. 5652138th Torrey Pines Road, Suite 220,  
;; CITY: La Jolla  
;; STATE: CA  
;; COUNTRY: USA  
;; ZIP: 92037  
;; COMPUTER READABLE FORM:  
;; MEDIUM TYPE: Floppy disk  
;; COMPUTER: IBM PC compatible  
;; OPERATING SYSTEM: PC-DOS/MS-DOS  
;; SOFTWARE: Patentin Release #1.0, Version #1.25  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: US/08/276,852  
;; FILING DATE: 18-JUL-1994  
;; CLASSIFICATION: 514  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 08/178,302  
;; FILING DATE: 30-SEP-1993  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 07/954,148  
;; FILING DATE: 30-SEP-1992  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Fitting, Thomas  
;; REGISTRATION NUMBER: 34,163  
;; REFERENCE/DOCKET NUMBER: SCRI452P  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: 619-554-2937  
;; TELEFAX: 619-554-6312  
;; INFORMATION FOR SEQ ID NO: 156:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 13254 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: double  
;; TOPOLOGY: circular  
;; MOLECULE TYPE: DNA (genomic)  
;; US-08-276-852-156

Query Match 60.3%; Score 142.8; DB 1; Length 13254;  
Best Local Similarity 81.7%; Pred. No. 3.7e-34;  
Matches 165; Conservative 0; Mismatches 37; Indels 0; Gaps 0;  
QY 36 GCAGGCATCTAGTGTGTCGAGGACGTGACAGCTAGCCCGGGCTCGAGATCTGGGATCTG 95  
Db 8076 GCATGATCTCAATTAGTCAGCAACAGGCTCCCGCAGCAGCAAGTATGCAAGCATG 8135  
QY 96 CATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCGGCCCATCCGCCCTAACT 155  
Db 8136 CATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCGGCCCATCCGCCCTAACT 8195  
QY 156 CGCCCGAGTTCGGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTATGAGAG 215  
Db 8196 CGCCCGAGTTCGGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTATGAGAG 8255  
QY 216 GCCGAGGCGGCTCGGCTCTG 237  
Db 8256 GCCGAGGCGGCTCGGCTCTG 8277

RESULT 11  
US-08-276-852-170/c  
; Sequence 170, Application US/08276852  
; Patent No. 5652138  
; GENERAL INFORMATION:  
; APPLICANT: Burton, Dennis R  
; APPLICANT: Barbas, Carlos F  
; APPLICANT: Lerner, Richard A  
; TITLE OF INVENTION: HUMAN NEUTRALIZING MONOCLONAL ANTIBODIES

;; TITLE OF INVENTION: TO HUMAN IMMUNODEFICIENCY VIRUS  
;; NUMBER OF SEQUENCES: 170  
;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: The Scripps Research Institute, Office of  
;; ADDRESSEE: Patent Counsel  
;; STREET: 10666 No. 5652138th Torrey Pines Road, Suite 220,  
;; CITY: La Jolla  
;; STATE: CA  
;; COUNTRY: USA  
;; ZIP: 92037  
;; COMPUTER READABLE FORM:  
;; MEDIUM TYPE: Floppy disk  
;; COMPUTER: IBM PC compatible  
;; OPERATING SYSTEM: PC-DOS/MS-DOS  
;; SOFTWARE: Patentin Release #1.0, Version #1.25  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: US/08/276,852  
;; FILING DATE: 18-JUL-1994  
;; CLASSIFICATION: 514  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 08/178,302  
;; FILING DATE: 30-SEP-1993  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 07/954,148  
;; FILING DATE: 30-SEP-1992  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Fitting, Thomas  
;; REGISTRATION NUMBER: 34,163  
;; REFERENCE/DOCKET NUMBER: SCRI452P  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: 619-554-2937  
;; TELEFAX: 619-554-6312  
;; INFORMATION FOR SEQ ID NO: 170:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 13254 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: double  
;; TOPOLOGY: circular  
;; MOLECULE TYPE: DNA (genomic)  
;; US-08-276-852-170

Query Match 60.3%; Score 142.8; DB 1; Length 13254;  
Best Local Similarity 81.7%; Pred. No. 3.7e-34;  
Matches 165; Conservative 0; Mismatches 37; Indels 0; Gaps 0;  
QY 36 GCAGGCATCTAGTGTGTCGAGGACGTGACAGCTAGCCCGGGCTCGAGATCTGGGATCTG 95  
Db 5179 GCATGATCTCAATTAGTCAGCAACAGGCTCCCGCAGCAGCAAGTATGCAAGCATG 5120  
QY 96 CATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCGGCCCATCCGCCCTAACT 155  
Db 5119 CATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCGGCCCATCCGCCCTAACT 5060  
QY 156 CGCCCGAGTTCGGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTATGAGAG 215  
Db 5059 CGCCCGAGTTCGGCCCATCTCCGCCCATCGCTGACTAATTTTTTTTATTATGAGAG 5000  
QY 216 GCCGAGGCGGCTCGGCTCTG 237  
Db 4999 GCCGAGGCGGCTCGGCTCTG 4978

RESULT 12  
US-08-899-575-156  
; Sequence 156, Application US/08899575  
; Patent No. 5770440  
; GENERAL INFORMATION:  
; APPLICANT: Burton, Dennis R  
; APPLICANT: Barbas, Carlos F  
; APPLICANT: Lerner, Richard A  
; TITLE OF INVENTION: HUMAN NEUTRALIZING MONOCLONAL ANTIBODIES  
; TITLE OF INVENTION: TO HUMAN IMMUNODEFICIENCY VIRUS

TITLE OF INVENTION: TO HUMAN IMMUNODEFICIENCY VIRUS  
NUMBER OF SEQUENCES: 170  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: The Scripps Research Institute, Office of  
ADDRESS: Patent Counsel  
SYNOPSIS: 10666 No. 5770440th Torrey Pines Road, Suite 220,  
STREET: Mail Drop TPC8  
CITY: La Jolla  
STATE: CA  
COUNTRY: USA  
ZIP: 92037

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41F: 2007
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
COMMAND: DATEINT Release #1.0. Version #1.25

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SOFTWARE: PatentIn Release #1.0.7  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/899,575  
FILING DATE: 24-JUL-1997  
CLASSIFICATION: 435

CLASSIFICATION: UNCLASSIFIED  
PRIORITY: UNCLASSIFIED  
PRIORITY APPLICATION NUMBER: US 08/276,852  
FILING DATE: 18-JUL-1994  
APPLICATION NUMBER: US 08/178,302

FILING DATE: 30-SEP-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/954,148  
FILING DATE: 30-SEP-1992  
AGENT: TROMBAY, JAMES

ATTORNEY/AGENT INFORMATION:  
NAME: Fitting, Thomas  
REGISTRATION NUMBER: 34,163  
REFERENCE/DOCKET NUMBER: SCRI452P  
TELECOMMUNICATION INFORMATION:

TELEPHONE: 619-554-2937  
TELEFAX: 619-554-6312  
INFORMATION FOR SEQ ID NO: 170:  
SEQUENCE CHARACTERISTICS:

LENGTH: 13254 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: circular

MOLECULE TYPE: DNA (genomic)	Score
8-899-575-170	60.38
very match	91.79

36 GCAGGCATCTAGTGTCTGTCGAGGAGCTGAC/

5179 GCATGCATCTCAATFAGTCAGCAACACGAGC  
96 CATCTCAATAGTCAGCAACCATAGTCCCG

5119 CATCTCAATTAGTCAGCAACCATAGTCCCGG  
156 CCGCCCCAGTTCGGGCCATTTCTCGGCCCAT

5059 CGGCCCAGTTCGGGCCAATTCTCGGCCCAAT  
216 GCGAGGCGGCTCGGCCTCTG 237  
|||||  
4000 CGGACCGGCGCTCGGCCTCTG 4978

4999 GCGAGGCGCCCTCGGCCCTC 4970

ULT 14  
08-899-575-156

sequence 156, Application US/08899575  
 Patent No. 5804440  
 GENERAL INFORMATION:  
 APPLICANT: Burton, Dennis R

APPLICANT: Barbas, Carlos F  
APPLICANT: Lerner, Richard A

NUMBER OF SEQUENCES: 170  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: The Scripps Research Institute, Office of  
ADDRESSEE: Patent Counsel  
STREET: 10666 No. 5770440th Torrey Pines Road, Suite 220,  
STREET: Mail Drop TPC8  
CITY: La Jolla  
STATE: CA  
COUNTRY: USA  
ZIP: 92037

ZIP: 32037  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 COMMUN.PP. Patent'n Release #1.0.  
 Version #1.25

SOFTWARE: PatentIn Release #1.0,  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/899,575  
FILING DATE: 24-JUL-1997  
CLASSIFICATION: 435

CLASSIFICATION:   
PRIOR APPLICATION DATA:   
APPLICATION NUMBER: US 08/276,852   
FILING DATE: 18-JUL-1994   
APPLICATION NUMBER: US 08/178,302

FILING DATE: 30-SEP-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/954,148  
FILING DATE: 30-SEP-1992  
----- AGENT INFORMATION:-----

ATTORNEY/AGENT INFORMATION:  
NAME: Fitting, Thomas  
REGISTRATION NUMBER: 34,163  
REFERENCE/DOCKET NUMBER: SCRL452P  
TELECOMMUNICATION INFORMATION:

TELEPHONE: 619-554-2937  
TELEFAX: 619-554-6312  
INFORMATION FOR SEQ ID NO: 156:  
SEQUENCE CHARACTERISTICS:

LENGTH: 13254 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: circular

MOLECULE TYPE: DNA (genomic)  
 SS-08-899-575-156

Query Match	60.3%	Score
Percent Similarity	81.7%	Pred

Best Local Similarity 81.1%; Predicted: 0; Mismatches 165; Conservative 0; Missed 0

db 8076 GCATGCATCTCAATTAGTCAGCAACCAGGC  
2y 96 CATCTCAATTAGTCAGCAACCATAGTCCCG

8136 CATCTCAATTAGTCAGCAACCATAGTCCCCG  
156 CCGCCCAGTTCGGCCCATTTCTCCGCCCCCAT  
|||||

db	8196	CCGCCAGTTCGCCCATTTCTCGGCCCA
QY	216	GCCGAGGCGCCTCGGCCTCTG 237
	8256	CGCCAGGCGCCTCGGCCTCTG 8277

Db 8256 GCGAGGCGCGCCCGCCCGCCCG 8277

RESULT 13

US-08-899-575-170/c

US 08 933 375 1787C  
; Sequence 170, Application US/08899575  
; Patent No. 5770440  
; GENERAL INFORMATION:  
; APPLICANT: Burton, Dennis R

APPLICANT: Barbas, Carlos F  
APPLICANT: Lerner, Richard A  
TITLE OF INVENTION: HUMAN NEUTRAL